

AP20 Rec'd PCTO 15 JUN 2006

**SYSTEMS FOR PREPARING FINE PARTICLES AND OTHER SUBSTANCES****FIELD OF INVENTION**

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This invention relates to controlled preparation of fine particles such as nano-crystalline films and powders with at least one solvent being in a supercritical state. It provides methods, measures, apparatus and products produced by the methods. In other aspects, the invention relates to further treatment of formed particles such as encapsulation of  
10 formed primary particles, and methods and measures for collection of formed substances in a batch wise, semi-continuous or continuous manner.

**BACKGROUND**

15 There is an increasing interest in nano- and micron sized materials in numerous technical applications. Such nanostructured fine particle materials in the form of nanocrystalline films and powders are cornerstones in the attempt to develop and exploit nanotechnology. They exhibit properties, which are significantly different from those of the same materials of larger size. During the last decade, the insight into nanostructured materials have  
20 dramatically improved through the application of new experimental methods for characterization of materials at the nanoscale. This has resulted in the synthesis of unique new materials with unprecedented functional properties. For nanostructured coatings, physical properties such as elastic modulus, strength, hardness, ductility, diffusivity, and thermal expansion coefficient can be manipulated based on nanometer control of the  
25 primary particle or grain size. For nano structured powders parameters such as the surface area, solubility, electronic structure and thermal conductivity are uniquely size dependent.

The novel properties of such nanostructured materials can be exploited and numerous new applications can be developed by using them in different industries. Examples of potential  
30 applications include new materials such as improved thermoelectric materials, electronics, coatings, semiconductors, high temperature superconductors, optical fibres, optical barriers, photographic materials, organic crystals, magnetic materials, shape changing alloys, polymers, conducting polymers, ceramics, catalysts, electronics, paints, coatings, lubricants, pesticides, thin films, composite materials, foods, food additives, antimicrobials,  
35 sunscreens, solar cells, cosmetics, drug delivery systems for controlled release and targeting, etc.

Addressing and exploiting such promising applications with new materials generally requires an improved price-performance ratio for the production of such nanostructured

materials. The key parameters determining the performance are the primary particle (grain) size, size distribution of the primary particles, chemical composition and chemical purity as well as the surface area of powders, while the primary parameters for In relation to price are the ease of processing and suitability for mass production.

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Several techniques have been used in the past for the manufacture of micron- or nano sized particles. Conventional techniques for submicron powders include spray drying, freeze drying, milling and fluid grinding, which are capable of producing powders in the micrometer range. Manufacturing techniques for producing submicron materials include  
10 high temperature vapour phase techniques such as flame synthesis and plasma arc methods, which allow production of nano-scaled powders consisting of hard or soft agglomerates of primary particles.

Solution sol-gel and hydrothermal synthesis are the major low temperature processes for  
15 production of fine particles with nano-scaled primary particles or grains. Hydrothermal synthesis is used for synthesis of a wide range fine oxide powders. The term hydrothermal relates to the use of water as reaction medium and regime of high pressure and the medium to high temperature applied. A major drawback is the relatively long reaction time required at for at low to medium temperatures and the very corrosive environment at  
20 higher temperature.

Sol-gel processing is widely used as it is a versatile technology that allows production of homogeneous high purity fine particles with a relatively small primary particle size to be produced from numerous materials in the form of powders, films, fibres, spheres,  
25 monoliths, aerogels, xerogels as well as coatings. The precursors can be metal organics, metals, inorganic salts etc. The processing temperatures are generally lower than for hydrothermal synthesis.

The key drawbacks from the sol-gel process are that it is that it is time consuming, and  
30 need after treatment such as drying and calcinations. In the traditional sol-gel process, it is necessary to calcine the product for up to 24 hours in order to obtain a crystalline product. In addition to a higher energy usage and a more complicated process this has the unfortunate effect that substantially growth of primary particles occur, and that the specific surface area may be decreased by up to 80 %.

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#### *Supercritical fluids*

Supercritical fluids exhibits particular attractive properties such as gas-like mass transfer properties like diffusivity, viscosity, and surface tension, yet having liquid-like properties

such as high solvation capability and density. Furthermore, the solubility can be manipulated by simple means such as pressure and temperature. This tunable solvation capability is a unique property that make supercritical fluids different from conventional solvents. Another major advantage of supercritical fluids is that rapid separation of solutes  
5 can easily be achieved by reduction of pressure. These attractive properties of such fluids at supercritical conditions have attracted considerable attention for its potential applications as environmentally friendly solvents for chemical processing. Carbon dioxide is the most widely used fluid for dense fluid applications, because of its moderate critical constants ( $T_c=31,1$  C,  $P_c=72,8$  atm, and  $\rho_c = 0,47$  g/cm<sup>3</sup>), non-toxic nature, low cost, and  
10 availability in pure form.

Supercritical CO<sub>2</sub> are today a mature technology, which are commercially being applied in large scale for extraction applications such as decaffeination of coffee and tea, extraction of hops, spices, herbs and other natural products. More recently supercritical fluids such as  
15 supercritical CO<sub>2</sub>, have been applied for commercial applications within impregnation.

Production of micron and submicron sized powders by supercritical techniques have been a hot scientific topic since the beginning of the nineties. The development has particularly been focused on physical transformation processes. They are generally variations of two  
20 primary methods for particle precipitation in supercritical fluids, the Solvent-AntiSolvent technique (SAS) and the Rapid Expansion of Supercritical Solutions technique (RESS).

#### SAS Technique

In the SAS technique, the material of interest is first dissolved in a suitable organic  
25 solvent, and the solution is subsequently mixed with a supercritical solvent, which dissolves the solvent and precipitates the solids out as fine particles.

#### RESS Technique

In the RESS technique, the solid of interest is first dissolved in a supercritical fluid and  
30 thereafter expanded by spraying through a nozzle. The expansion through the nozzle causes a dramatic reduction in the CO<sub>2</sub> density and thereby a dramatic reduction in the solvent capacity, causing high supersaturation resulting in the formation of fine particles.

Derived techniques from the SAS and RESS techniques are for example Solution Enhanced  
35 Dispersion by Supercritical Fluids Techniques (SEDS) and Precipitation with compressed Antisolvent technique (PCA), which is based on the concept of coupling the use of a supercritical fluid as a dispersing agent, by means of a coaxial nozzle, in addition to its primary role as an antisolvent and a vehicle to extract the solvent. Further extensions of this technique include multiple concentric opening nozzles.

Other techniques include Precipitation from Gas-Saturated Solutions (PGSS), which involves melting the material to be processed, and subsequently dissolving a supercritical fluid under pressure. The saturated solution is then expanded across a nozzle, where the  
5 more volatile supercritical fluid escapes leaving dry fine particles.

All these techniques have been successfully used in small scale to produce micron sized particles of various materials for numerous applications. Excellent reviews of prior art supercritical particle formation processes can be found in e.g. Ya-Ping Sun ("Supercritical  
10 Fluid Technology in Materials Science and Engineering – Syntheses, Properties and Applications, Marcel Dekker Inc., 2002-ISBN: 0-8247-0651-X), Gentile et al (WO03/035673A1), Gupta et al (US2002/0000681A1), Mazen et al (EP0706421B1), Del Re et al (WO02/068107A2), Mazen et al (WO99/44733), Calfors et al, Jagannathan et al (WO03/053561), all of which are hereby included by reference.

15 However, all these techniques suffer from some inherent limitations. The RESS technique is limited by the solvent capacity in the supercritical fluid. For example, supercritical carbon dioxide, which is a preferred solvent in many applications, is limited by a low solubility towards polar substances. Modifiers such as co-solvents and surfactants may be  
20 added to the supercritical carbon dioxide to improve the solubility of the material of interest. However, such co-solvents and surfactants may remain in the precipitated product as impurities, which may not be acceptable. Further drawbacks of the RESS technique includes that the isenthalpic expansion over the nozzle that results in large temperature drops, which can cause freezing of the solid and carbon dioxide and thereby  
25 cause blocking of the nozzle. The nozzle design is further critical for the final particle characteristics such as size and morphology etc. All these drawbacks from microscopic variables limit the control over the process itself, and make scale-up relatively difficult. Still further such systems are in its present embodiment generally limited to non-reacting or extremely fast reacting systems as the change of solubility is caused momentary.

30 Due to the higher solubility the SAS technique and its derivatives generally have higher through-puts, and generally produce particles in the range 1-10 micron (Gupta et al, US2002/0000681A1). The key and particle size controlling step of the SAS techniques is the mass transfer rate of the antisolvent into the droplet. Hence, mixing of solution and  
35 the supercritical fluid is crucial In order to obtain an intimate and rapid mixing, a dispersion of solution as small droplets into the supercritical fluid is required. Various nozzle designs have been proposed to inject solution and supercritical fluid into a particle formation vessel in order to provide a good mixing. Recent modifications of the SAS technique to reduce the particle size includes atomization techniques such as special designed coaxial nozzles,

vibrational atomization, atomization by high frequency sound waves, ultrasonic atomization etc. (US2002000068A1). Though these modified techniques are believed to provide enhanced mass transfer and resulting reduced particle sizes, too rapid particle formation may reduce the control of the size and morphology such as crystallinity of the formed particles, be sensitive to the nozzle design and blockages of the nozzle and be difficult to scale-up. A further drawback is that the SAS techniques are generally not suitable for reactive systems in large scale.

## DESCRIPTION OF THE INVENTION

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A major shortcoming in the widespread commercial exploitation of nanotechnology has so far been large scale production of fine particles with sufficient homogeneity and reproducibility at affordable costs so as to make them competitive in the market.

15 Fine particles in the present context generally comprise primary particles such as grains, crystallites and the like. It should be understood that the fine particles in this context, shall preferably be interpreted in broad terms. Said fine particles may comprise anything from a single primary particle, a cluster or clusters of primary particles, agglomerates of primary particles such as a powder, a film or a coating of said primary particles or even a bulk material comprised by said primary particles.

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Different aspects of the present invention seek to meet one or more of the following objectives:

25 An objective of present invention is to address the quality and availability of such fine particles by providing method(s) for production of such materials, which allows production of more homogeneous fine particles than in the prior art i.e. fine particles with a high purity and/or a controlled particle morphology, and/or a small average diameter and/or a narrow size distribution, and/or a controlled phase and/or structure.

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Another objective of the present invention is to provide method(s), which allow such high quality materials to be produced at shorter processing times and/or at lower temperatures and/or with a more controlled growth rate and/or with a more controlled morphology such as a more controllable crystallinity or shape than hitherto.

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Still another objective of the present invention is to provide method(s) suitable for large scale production of fine particles with more uniform and/or homogeneous properties.



A further objective of the present invention is to provide improved methods and measures for introducing fluid(s), and/or chemical reactant(s) and/or Initiator(s) and/or precursor(s) and/or catalyst(s) into a vessel.

- 5 A still further objective of the present invention is to provide improved methods and measures for controlling a chemical reaction in a dense fluid under near or supercritical conditions.

Still another further objective of the present invention is to provide methods which reduce  
10 or eliminates the needs for post processing steps such as drying and calcinations.

Furthermore, an objective of the present invention is to provide methods and measures for the collection the fine particles in both a batch wise manner and a continuous manner.

- 15 It may also be an objective of the present invention to provide an apparatus for production of fine particles according to the above described method.

Additionally, it may be an objective to provide a product obtained by the above described methods, and applications for use of said product.

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These objectives and the advantages that will be evident from the following description are obtained by the following preferred embodiments of the invention.

In a first aspect, the present invention of relates to the production of fine particles. Hence,  
25 a preferred embodiment of a method according to the present invention comprises producing a fine particle material by

- i) introducing one or more substances contained, such as dissolved and/or dispersed  
in one or more fluid(s) into a vessel by introducing said fluid(s) into the vessel, said  
30 vessel containing one or more section(s) comprising a material, at least one of the  
fluids being in a supercritical state before or after being introduced into said vessel,  
ii) causing and/or allowing said substances to precipitate at least partly as primary  
particles on the surface of said material.

35 In many embodiments according to the present invention, the method relates to the production of fine particles comprising nanoscaled primary particles i.e. primary particles having an average diameter smaller than 100 nanometer such as smaller than 30 nanometer, and even more preferable primary particles having an average diameter of smaller than 15 nanometer such as an average diameter smaller than 10 nanometer.

As the primary particles may have an irregular shape, the average diameter in this context shall preferably be interpreted as an equivalent spherical diameter. Various methods of varying quality exist for the determination of the size of nanoscaled primary particles including X-Ray Diffraction (XRD), Small Angle X-ray Scattering (SAXS), Transmission Electron Microscopy (TEM), Scanning Electron Microscopy (SEM). The microscopic techniques may lead to inaccuracies, and it is recommended to apply the X-ray techniques. The above average diameters refers to diameters determined by the SAXS technique by applying the Beaucage model [G. Beaucage et al, Journal of Non-crystalline Solids 172-174, p. 797-805, 1994]. This method is considered as reliable and widely applicably as it allows determination of the average diameter of both amorphous and crystalline phases.

Many preferred embodiments according to the present invention relates to the production of very uniform and homogeneous fine particle materials having a very narrow size distribution.

Hence, a method according to the present invention often comprises fine particles, wherein the standard deviation of the size distribution of the average diameter of said primary particles formed is often less than 60 % of the average diameter, such as 40 % of the average size of said primary particles, and preferably less than 30 % of the average size of said primary particles such as less than 20 % of the average size of said primary particles, and even more preferably the standard deviation of the size distribution of said primary particles formed is less than 15 % of the average diameter of said primary particles.

A preferred embodiment of the present invention relates to the production of fine particles wherein the standard deviation of the average diameter of said primary particles formed is maximum 20 nanometer, such as maximum 10 nanometer, and preferably less than 5 nanometer, and even more preferably less than 3 nanometer. The above mentioned standard deviation may be derived from SAXS data or similar high quality data.

The present invention generally relates to a method, wherein at least one of said fluids is/are in a supercritical state before or after being introduced into said vessel. In a preferred embodiment said fluid(s) being in a supercritical state is/are preferably selected from the group consisting of carbon dioxide, alcohols such as methanol, ethanol, propanol, isopropanol, butanol, sec-butanol, pentanol, hexanol, water, methane, ethane, propane, butane, pentane, hexane, cyclohexane, heptane, ammonia, sulfurhexafluoride, nitrous oxide, chlorotrifluoromethane, monofluoromethane, acetone, THF, acetic acid, citric acid, ethylene glycol, polyethylene glycol, N,N-dimethylaniline and mixtures thereof.

In particular preferred embodiments at least one of the fluid(s) may be CO<sub>2</sub> and/or an organic solvent and/or water.

In another embodiment said fluid may further comprise at least one co-solvent preferably  
5 selected from the group consisting of alcohol(s), water, ethane, ethylene, propane, butane, pentane, hexane, heptane, ammonia, sulfurhexafluoride, nitrous oxide, chlorotrifluoromethane, monofluoromethane, methanol, ethanol, propanol, isopropanol, butanol, pentanol, hexanol, acetone, DMSO, THF, acetic acid, ethyleneglycol, polyethyleneglycol, N,N-dimethylaniline and mixtures thereof.

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In yet another embodiment said fluid may also comprise one or more surfactants, said surfactants being preferably selected from the group consisting of hydrocarbons and fluorocarbons preferably having a hydrophilic/lipophilic balance value of less than 15, where the HLB value is determined according to the following formula:  $HLB = 7 +$   
15  $\text{sum}(\text{hydrophilic group numbers}) - \text{sum}(\text{lipophilic group numbers})$ .

The pressure of at least one of said fluids being in a supercritical state before or after being introduced to the vessel may be in the range 85-500 bar, preferably in the range 85-350 bar, such as in the range 100-300 bar. In embodiments, wherein said fluid(s) being in  
20 a supercritical state before being introduced to the vessel and not within the vessel said fluid(s) often undergo an expansion into said vessel according to methods well known in the prior art.

However, many preferred embodiments of the present invention relates to methods,  
25 wherein the at least one of said fluid(s) being introduced into said vessel, is in an supercritical state also after introduction into said vessel. In such embodiments the pressure within the vessel may be in the range 85-500 bar, preferably 85-350 bar such as in the range 100-300 bar.

30 The absolute temperature depends of the actual fine particles to be produced and may in many embodiments according to the present invention be maintained in the range 20-500 °C, such as 30-450 °C, and preferable in the range 35-200 °C, and more preferable in the range 40-150 °C.

35 The precipitation is generally caused by change of the solubility of at least one of said substances. The change of said solubility may be performed in a number of ways depending of the specific particle formation application.



In one embodiment, said changing of the solubility involves mixing said fluid(s) containing said dissolved and/or dispersed substances with an antisolvent capable of dissolving at least partly at least one of said fluid(s) and/or a reaction product formed by a chemical reaction occurring as a result of said mixing. The antisolvent may be in a gaseous, liquid or  
5 a supercritical state. The antisolvent may be present in the vessel prior to introducing said fluid(s) and/or may be introduced into said vessel together with said high surface area material at one or more points.

In another embodiment said changing of the solubility of at least one of said substances,  
10 may be to expand the fluid(s) containing said substances into the vessel through one or more nozzles such as performed in the Rapid Expansion of Supercritical Solvent (RESS) and the Rapid Expansion of Supercritical solvent into a Liquid (RESOLV) techniques. Still another embodiment involves changing the solubility by changing the temperature of said fluid.

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In a preferred embodiment according to the present invention, at least one of said dissolved and/or dispersed substances in said fluid(s) undergoes a chemical reaction. Said reaction may be a reaction according to the so-called sol-gel route. Traditional sol-gel processing is versatile and widely used reaction route, which allows synthesis of a wide  
20 range of materials including oxides, hydroxides, oxyhydroxides, nitrides, carbides etc. of e.g. metals or semi-metals.

A good description of the traditional sol-gel synthesis method for e.g. making fine ceramic fibers as described in e.g. YA-Ping Sun, "Supercritical Fluid Technology in Materials Science  
25 and Engineering – Syntheses, Properties, and Applications", Marcel Dekker, 2002, ISBN:0-8247-0651-X hereby included by reference. It involves forming an aqueous dispersion of oxide particles that is then gelled either by concentrating the dispersion by solvent removal or by carrying out a chemical reaction. For example, one method of sol gel synthesis is to start with a metal alkoxide solution and add a small amount of water to control the  
30 hydrolysis and condensation of metal hydroxides. As the sol is dried, these metal hydroxides form a polymeric network through cross linking of the metal oxygen bonds. The method of drying greatly influences the final product morphology. Supercritical drying has been shown to produce soft aggregates that can be broken down to a powder. The resulting powder is typically subjected to heat treatment to induce the complete  
35 dehydration and crystallization of oxide particles. Another method of sol-gel synthesis is to start with a solution of a metal salt and a water-soluble polymer. By adding a base to this solution, the metal salt can be converted to metal hydroxides, while the polymer cross-links to form a porous network around these metal hydroxides. In this case, the polymer network serves to prevent significant growth and aggregation of the metal hydroxides, so

that it is possible to obtain nanomaterials using this technique. Nanomaterials of a number of ferrites including  $\text{CoFe}_2\text{O}_4$ ,  $\text{Ni}_{0,5}\text{Zn}_{0,5}\text{Fe}_2\text{O}_4$ ,  $\text{BaFe}_{12}\text{O}_{19}$  and  $\text{Ge}_{0,5}\text{Fe}_{2,5}\text{O}_y$  have been prepared using sol-gel synthesis.

- 5 In an embodiment of the present invention a sol-gel reaction may be performed with an alkoxide precursor dissolved in e.g. supercritical  $\text{CO}_2$  and/or an alcohol such as ethanol, isopropanol, butanol and/or super. The metal alkoxide reacts readily with water to produce metal oxides and/or metal hydroxides. Compared with samples made via conventional sol-gel syntheses, supercritical synthesized powders exhibit a higher degree  
10 of crystallinity and contains less hydroxide.

A particular advantageous embodiment of present invention leading to improved control of the properties of the primary particles formed may involve introducing the reactants sequentially. In such embodiments according to the present invention, it is advantageous  
15 to introduce at least one reactant(s) and/or precursor and/or initiator(s) and/or catalyst(s) into the vessel at least partly prior to introducing said fluid(s) containing said substances and vice versa.

Furthermore, in an embodiment according to the present invention such sequential or  
20 stepwise introduction of reactants may be repeated multiple times e.g. by introducing into the vessel at least one of said reactant(s) and/or precursor(s) and/or initiator(s) for said chemical reaction and subsequently introducing into the vessel one or more substances dissolved and/or dispersed in or mixed with at least one fluid or vice versa.

25 Additionally, such multiple sequential steps may also involve one or more of the following processes: RESS (rapid expansion of supercritical solutions), GAS (Gas Antisolvent), SAS (solvent Anti Solvent), SEDS (Solution Enhanced Dispersion by supercritical fluid), PCA (Precipitation with Compressed Antisolvent), PGSS (Precipitation from Gas-saturated Solutions) and variations thereof either prior to introducing into the vessel at least one of  
30 said reactant(s) and/or precursor(s) and/or initiator(s) for said chemical reaction and subsequently introducing into the vessel one or more substances dissolved and/or dispersed in or mixed with at least one fluid, or after one or more of such sequential steps.

It may further be advantageous, if said material is capable of adsorbing at least one of  
35 said reactant(s) and/or precursor(s) and/or initiators(s) and/or catalyst(s), and preferably in substantially a monolayer. Hereby said reactant can be evenly distributed on said high surface area material thereby resulting in a very controllable reaction and/or fine particle formation process.

The temperature in the vessel may be selected so as to control the specific properties of the primary particles formed e.g. crystallinity, particles size and phase. A higher temperature generally leads to higher reaction rates, but also reduces the control of the specific properties. A particular feature of the present invention may be that it allows  
5 controlled production of homogeneous materials at higher reaction rates and lower temperatures than hitherto.

As described above the temperature in the vessel during said sol-gel reaction generally depend on the specific fine particle material to be produced. In many preferred  
10 embodiments the maximum temperature in the vessel during said sol-gel reaction(s) may be maintained below 400 C, such as below 300 C, preferably below 250 C such as below 200 C, and even more preferably below 150 C such as below 100 C.

The temperature may further be maintained constant during each of said sequential steps,  
15 or may be varied according to a pre-selected schedule. In embodiments, wherein different materials are produced in the individual steps the temperature and/or pressure may further be varied between each of such individual steps.

In some embodiments according to the present invention the time for said chemical  
20 reaction may be relatively long such as less than 24 hours, such as less than 12 hours, and preferable less than 8 hours such as less than 4 hours.

In a preferred embodiment according to the present invention the time for said chemical  
reaction(s) is maximum 2 hours, such as maximum 1 hour, preferably less than 30  
25 minutes and even more preferably less than 15 minutes.

The material being present in said one or more sections according to many embodiments of the present invention may have a number of functions. In some embodiments according to the present invention, it may serve as a distributor enabling a more uniform distribution  
30 of the substances being introduced into the vessel, and thereby improve the homogeneity of the fine particle material(s) being formed.

In other embodiments to the present invention the material may provide a large number of nucleation sites so as to provide a high nucleation rate compared to the particle growth  
35 rate. Hereby, a seeding effect may be introduced, thus ensuring a fine control of the fine particles formed. In some embodiments the seeding effect may further be increased by introducing an ultrasound and/or a vibrating effect. In other embodiments said seeding effect may further be at least partly provided by seed particles.

The material may be arranged in a number of ways in said one or more section(s) of said vessel. The vessel may be whole or partly filled with said material. In many embodiments according to the present invention the material may comprise a porous structure and the fraction of the total volume comprised by said material in said one or more sections may  
5 be up to 70 %, such as up to 50 %, preferably up to 30 % and even more preferably up to 20 %.

In one embodiment according to the present invention said particles may be in a fluidised or suspended state in said one or more section in the vessel. In a further embodiment  
10 according to the present invention the material may comprise the same material as said primary particles.

The porous structure of said material in said one or more sections may have any shape, such as a sheet, a fibrous, a spongy or a grid structure. In a preferred embodiment  
15 according to the present invention said material present in said one or more sections may be a template for forming and/or curtailing said primary particles into a specific shape, size, structure or phase.

The material may comprise a wide range of materials depending on the specific  
20 application. In many embodiments according to the present invention the material may be selected so as to provide a specific functionality. One such functionality may be the capability to adsorb specific compounds on the surface, whereby specific properties of the formed fine particle product may be controlled e.g. average particle diameter and size distribution.

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In an embodiment according to the present invention involving water as a precursor/initiator for a reaction, the material may be selected so as to provide a large adsorption capacity for water. In such cases a hydrophilic material is selected.

30 In another embodiment it may be desired to obtain a selective adsorption of another substance e.g. an alkoxide. In such embodiments a less hydrophilic or a hydrophobic material may be selected.

In many embodiments according to the present invention said material may comprise a  
35 polymer material such as a polymer or elastomer selected from the group consisting of polyethylene, polypropylene, polystyrene, polyesters, polyethylene terephthalate, polyvinyl chloride, polyvinyl acetates, polyoxymethylene, polyacrylamide, polycarbonate, polyamides, polyurethane, copolymers thereof, chlorinated products thereof, rubbers and chlorinated rubber, silicone rubbers, butadiene rubbers, styrene-butadiene-rubbers, isoprene

polymers, vulcanised fluororubbers, silicone rubbers.

In a preferred embodiment according to the invention the polymer material may be polypropylene. In another preferred embodiment according to the present invention the material may comprise an elastic material. In still another preferred embodiment according to the present invention may be a ceramic material such as glass wool such quartz wool.

In a further embodiment according to the present invention the material may comprise a porous media such as an aerogel. In a particular preferred embodiment according to the present invention said aerogel may be produced within the same equipment by producing said aerogel material by

- a sol-gel reaction in an organic solvent
- removing said organic solvent by extraction in supercritical CO<sub>2</sub>
- drying at least partly said aerogel by supercritical CO<sub>2</sub>
- forming said primary particles on the surface of said aerogel according to the present invention.

In still another embodiment of the present invention said porous media material may comprise as a heterogeneous catalyst support material or a heterogeneous catalyst.

In many embodiments according to the present invention said material may comprise a high surface area material. Such high surface area materials according to the present invention may have a specific surface area (m<sup>2</sup>/m<sup>3</sup>) of said material in said sections is above 500 m<sup>2</sup>/m<sup>3</sup>, such as 1000 m<sup>2</sup>/m<sup>3</sup>, such as above 10.000 m<sup>2</sup>/m<sup>3</sup>, and preferably above 50.000 m<sup>2</sup>/m<sup>3</sup> such as above 100.000 m<sup>2</sup>/m<sup>3</sup>.

The high surface area material may comprise a plurality of fibres. Various ways of arranging such fibres are known in the prior art (e.g. W.S. Winston Ho et al et al, "Membrane Handbook", Van Nordstrand Reinhold, 1992, ISBN 0-442-23747-2, K. Scott, "Handbook of Industrial Membranes", Elsevier Science Publicers, 1995, ISBN 1856172333, Iversen et al, WO95351153, Iversen et al, WO00160095, US690,830, US5,690,823) and are hereby included by reference. Such methods includes random packings, mats, cloths , bundles, twisted bundles, meshes, arrays, etc..

In an embodiment of the present invention said fibres may comprise a plurality of fibres extending in substantially the same direction, such as in a filtration medium. One way of packing such fibres relevant to the present invention is disclosed in US 5,690,823 hereby included by reference.



In the present description with claims the term the term "hollow tubular member(s)" comprises hollow fibres, and other hollow tubular bodies having any cross section, e.g. a hollow tubular chamber. Likewise the term surface of a membrane and similar expressions are intended to mean at least part of a membrane surface.

5

In general it may be advantageous to introduce at least partly one of said fluids into said vessel through the walls of at least one hollow tubular member comprising an inner and an outer surface, and having at least one end communicating with the outside of said vessel. At least part of said hollow tubular member(s) comprising a membrane. Said membrane may comprise a so-called dense membrane. The term dense membrane is known by a man skilled in the art, and is intended to designate membranes having at least one layer being substantially nonporous i.e. having pores of substantially molecular dimensions.

15 In many embodiments according to the present invention, the membrane(s) is porous. In other applications such membranes are used for filtration of e.g. liquids (nanofiltration, ultrafiltration, microfiltration etc.), and have pores within the range 0,001-100 micron, such as pores in the range 0,01-10 micron, and preferably in the range 0,01-0,1 micron.

20 In some embodiments in accordance with the present invention such hollow tubular members may be used for introducing at least one of said fluid(s) into the vessel in a very uniform manner. In such embodiments the high surface area may comprise both hollow tubular member(s) and other high surface area materials such as a nonporous fibre material. The hollow tubular member(s) may also comprise several sets of hollow tubular member(s) for introducing different fluids to the vessel. Various examples integrating of said hollow tubular member(s) in the vessel are further illustrated in the figures.

In a particular preferred embodiment for many applications, the hollow tubular member(s) constitutes said high surface area material. In such embodiment a fluid and/or reactant(s) and/or initiator and/or precursor may be added to substantially the outer surface of said tubular member(s).

30 An important embodiment of the present invention may comprise re-circulating in at least part time of the method at least part of a fluid mixture present in the vessel, the re-circulating comprising:

- withdrawing from the vessel at least a part of a fluid from the vessel and feeding it to a re-circulation loop and subsequently feeding the fluid back to the vessel.

A preferred embodiment according to the present invention may further comprise the step of controlling the temperature of the fluid in the re-circulation loop by adding or extracting heat from said fluid in said re-circulation loop.

- 5 In another preferred embodiment one or more reactant may be added and/or extracted from the fluid in said re-circulation loop, hence allowing precise control of said reactant concentrations during said fine particle material reaction. Still another preferred embodiment may involve controlling the concentration of an alcohol, an alkoxide and/or water. A further preferred embodiment according to the present invention may comprise
- 10 controlling the temperature- and/or pressure- and/or density- and/or concentration profiles within the vessel. Such embodiment increases the mass- and an heat transfer within the vessel and allow precise control of the fine particle material being formed.

In a particular preferred embodiment a metal- or semi metal alkoxide are produced in-situ

15 in the recirculation loop e.g. by applying an electrochemical synthesis of said metal or semi-metal in the corresponding alcohol. Said metal or semi-metal alkoxide may be introduced into said vessel in various ways such as exemplified in the figures 7-8, and in illustrative example 1.

- 20 An important embodiment according to the present invention may be where said material within said one or more sections in vessel with said precipitated primary particles thereon comprises the final product. Non-limiting examples of such product according to such embodiment comprises a tape cast with said primary particles deposited on a carrier film. Alternatively, wherein said primary particles on said surface of said material constitutes a
- 25 film or a coating.

A particular preferred embodiment according to the present invention may be wherein said film or coating has one or more layer(s) each layer having a layer thickness of up to 1 micron, such as a layer thickness below 500 nanometer, preferable a layer thickness below

30 250 nanometer such as a layer thickness below 100 nanometer. Even more preferable the layer thickness of said film is below 50 nanometer, such as a layer thickness below 30 nanometer. Additionally, said coating or film on said material may comprise multiple layers, and optionally these layers may comprise different materials.

- 35 Another embodiment according to the present invention may comprise subjecting said coating or film to an annealing process. In a preferred embodiment said annealing process may be performed by microwaves. Such has many advantages compared to conventional thermal annealing as it is clean and simple, energy and cost efficient and has a short reaction time. It may further be integrated in the production process. Another distinct

advantage is that it may be applied for annealing coating or films on materials such as glass and polymers, where the conventional thermal annealing process is a limiting factor.

A further embodiment according to the present invention may be related to deposition of  
5 said primary particles on the surface of said material in the form of small clusters of individual particles and preferably as individual particles. This has become possible due to the elimination of the normal drying phenomena related to wet deposition methods, by applying a highly tuneable supercritical process according to the present invention. In an preferred embodiment according to the present invention said clusters may comprise up to  
10 100 atoms, such as up to 50 atoms, and preferably less than 10 atoms and even more preferably less than 5 atoms. A particular preferred embodiment said clusters and/or individual particles may be deposited as quantum dots.

A further embodiment according to the present Invention may be where said primary  
15 particles precipitated on said surface of the material present in said one or more section(s) are removed from said material as a powder. In most embodiment according to the present invention said powder consists of weakly bounded soft agglomerates of primary particles. In many of such embodiments according to the present Invention said soft weakly bounded agglomerates may have a size of maximum 10 micron, such as up to 5  
20 micron, and preferably up to 1 micron such as up to 500 nanometer.

In a preferred embodiment according to the present invention said powder may be removed from said material by introducing a vibrating effect and/or an acoustic effect such as ultrasound waves and/or by back flushing and/or by applying a pressure pulse effect.  
25 The vibrating effect may in an embodiment according to the present Invention be generated by piezoelectric means.

Alternatively said vibrating effect may be generated by a magneto-restrictive means in accordance with an embodiment of the present invention.  
30

In a particularly preferred embodiment according to the present invention said material may be removed within the vessel, thus allowing for continuous or semi-continuous operation.

35 The removal of said removed powder from said material may be withdrawn from the vessel by flushing with a fluid or fluid mixture present in the vessel in accordance with an embodiment of the present invention.

Additionally, said fluid containing said formed powder may be fed into a second vessel containing a liquid in accordance with the present invention, and thereby providing said powder material as a dispersion in said liquid.

- 5 Alternatively, said fluid containing said powder may be fed to a bag filter or a ceramic filter for separation of said formed powder material from said fluid. Furthermore, said formed particulate material may be fed to a membrane separation device.

- 10 In another preferred embodiment according to the present invention, a coating or encapsulation step may be performed. In a particular preferred embodiment said coating or encapsulation step(s) may be performed at least partly during harvesting/removing said particles from said material.

- The present invention is applicable for production of fine particles from a wide range of materials. Preferred embodiments according to the present invention include the production of primary particles wherein said primary particles comprises oxide(s), oxyhydroxide(s), hydroxide(s) such as metal oxide(s), semi-metal oxide(s), metal oxyhydroxide(s), semi-metal oxyhydroxide(s), metal hydroxide(s), semi-metal hydroxides and combinations thereof.

20

Such preferred embodiments according to the present invention further include oxide materials such as electro-ceramic materials, semi-conducting materials, piezoelectric materials, and magnetic, ferromagnetic, paramagnetic, or super-paramagnetic materials.

- 25 In particular preferred embodiments according to the present invention said oxide materials may comprise oxides of one or more of the following elements: Al, Si, Ti, Zr, Zn, Fe, Ni, Co, Ce, Ge, Ba, Sr, W, La, Ta, Y, Mn, V, Bi, Sn, Te, Se, Ga, Be, Pb, Cr, Mg, Ca, Li, Ag, Au, Pt, Pd, Cd, Mo, Eu and combinations thereof.

- 30 In other embodiments according to the present invention said metal or semi-metal is/are precursors for a thermoelectric material. In an embodiment according to the present invention such materials are produced by applying a reducing agent to form a thermoelectric material.

- 35 In a preferred embodiment said thermoelectric material formed may comprise a clathrate, preferably comprising one or more of the following elements: Ba, Bi, Te, Se, Zn, Sn, Sr, Ga, Ge, Pb, Cd, Sb, Ag, Si and combinations thereof. An advantage of the present invention for such embodiments is that the small primary particles and the narrow size

distribution according to the present invention introduces an additional heat conductivity barrier between the primary particles.

Hence, in a preferred embodiment of the present invention a thermoelectric material  
5 having a thermal conductivity at temperatures above 20 C of maximum 10 watts per meter Kelvin, such as maximum 5 watts per meter Kelvin, preferably maximum 3 watts per meter Kelvin such as maximum 1,5 watts per meter Kelvin, and even more preferably a heat conductivity of maximum 1 watt per meter Kelvin may be produced. The primary particles of said thermoelectric material may further be doped with metals and/or semi-  
10 metals to improve the electrical conductivity of said material.

In second aspect of the present invention it further comprise an apparatus comprising one or more of the means disclosed in any of the preceding claims and being adapted to carry out the method according to the invention.

15

In a third aspect the present invention also relates to a product obtainable according to the invention.

A preferred embodiment according to the present invention may comprise a tape cast for  
20 tape casting, comprising primary particles deposited on a carrier film, wherein said primary particles have:

- a. an average diameter of less than 100 nanometer such as an average diameter of less than 30 nanometer, preferably an average diameter of  
25 smaller than 20 nanometer and even more preferable an average diameter below 15 nanometer such as below 10 nanometer.
- b. a narrow size distribution around the average diameter characterized by having a maximum standard deviation of said distribution of maximum 20 nanometer, such as maximum 10 nanometer, and preferably less than 5  
30 nanometer.

Another embodiment according to the present a piezomotor may be produced from a lead zirconate titanate tape cast..

35 An important embodiment according to the present invention may comprise an item having a hard nanocrystalline coating comprising primary particles of  $\text{Al}_2\text{O}_3$  and  $\text{ZrO}_2$  according to any of the preceding claims, wherein said coating has a hardness of at least 10 GPA, such as a hardness of at least 15 GPA, and preferably above 20 GPA, and even more preferably a hardness of at least 25 GPA.



A further embodiment according to the present invention may comprise an item having a hard nanocrystalline coating comprising primary particles of  $\text{Al}_2\text{O}_3$  and  $\text{ZrO}_2$  according to any of the preceding claims, wherein said coating has a scratch and wear resistance of at least 30 N, such as a scratch and wear resistance of at least 35 N, preferably a scratch and wear resistance of at least 40 N, and even more preferably a scratch resistance of at least 45 N.

It is well known in the prior art that a number of physico-chemical properties are uniquely size dependent, and that manipulation of such size dependent properties allow tailoring materials to specific applications.

The primary particles according to an embodiment to the present invention may be highly chemically pure and very homogeneous with a small and tunable average diameter, and a very narrow size distribution. Further said primary particles may be present in the form of a coating, a dry powder or in the form of a liquid suspension.

It will be known to a person skilled in the art that a number of applications for such products exist or may be developed including the applications mentioned in the described under background in this document.

## DESCRIPTION OF THE DRAWINGS

The following abbreviations applies to the figures below:

25

F: Fluid

F1: Fluid 1

F2: Fluid 2

HSAM: High surface area material

30 FP: Formed particles

PH: Particle harvesting

HTM: Hollow tubular member

Fig.1 shows an example of a vessel containing a high surface area fibre material according to the present invention. The high surface area material is contained in a vessel having one or more inlets for introducing one or more fluids. The vessel may be horizontally or vertically positioned. A randomly packed fibre material is illustrated in Fig. 1b. Fig. 1c shows a reactant (black triangles) adsorbed to said fibre material. Fig. 1d shows said

primary particles formed on the surface of said fibre surface, and Fig. 1e shows the harvesting said deposited particles

Fig.2 shows an example of a vessel similar to the one in Fig. 1, but further comprising a  
5 hollow tubular member blocked in one end to distribute said first fluid. It should be understood that the vessel may constitute a plurality of such tubular members.

Fig.3 shows a vessel containing a high surface area material according to the present invention comprising a plurality of fibres extending in substantially the same direction and  
10 with both ends communicating with the outside of said vessel. The vessel may have one or more inlets communicating with the outside of said vessel for introducing one or more fluids, and the vessel may further have one or more outlets for withdrawing said fluids and/or said particles formed. It should be understood that in addition to said high surface area material, the vessel may further comprise hollow tubular member(s) with one or both  
15 ends communicating with the outside of said vessel.

Fig.4. illustrates a vessel similar to the one in Fig. 3, but further comprising a plurality of hollow tubular members extending in substantially the same direction and communicating with both an inlet and an outlet plenum. The first fluid is introduced into said inlet plenum  
20 and is distributed to the inner surface of said tubular member(s). At least part of said fluid permeating through the membrane walls of said tubular members so as to obtain a controlled addition of said first fluid and/or dissolved substances to fluid on the outer surface of said hollow tubular members, thereby resulting in a precise control of the concentration of said fluid and/or dissolved substances within the vessel. The temperature  
25 within the vessel may further be precisely controlled by controlling the flow rate and inlet temperature of said first fluid. This is preferably accomplished by withdrawing in at least part of said particle formation process said first fluid from said outlet plenum to an external re-circulation loop (not shown), wherein flow rate, composition, temperature, and pressure are controlled in a predefined manner before re-circulating it to said inlet plenum  
30 for said first fluid. In a preferred embodiment the particles deposited on the outer surface of said hollow tubular members are at least partly removed from said surface by closing the outlet for said first fluid e.g. by closing a valve. Thereby substantially all of said first fluid permeates said membrane wall and clean the surface by back flushing. If said closing of the valve is very fast a back chock (short pressure pulse is obtained). It is further  
35 advantageous if said hollow tubular member is made from an elastic material so it is capable of expanding during said pressure pulse. It should be understood that the vessel may further comprise an additional high surface area material in addition to the hollow tubular members shown on the figure.

Fig. 5 shows an example of superimposed layers of hollow tubular members where two different fluids (A and B) can be conducted through the lumen of the fibres, as indicated, whereas a flow of a third fluid can be passed transversely through the fibres from above, perpendicular to the longitudinal direction of the fibres, as indicated by the vertical arrow.

5

Fig. 6 illustrates a situation similar to the one in Fig. 6, but where a woven array of hollow membrane fibres is used.

Fig. 7: illustrates a schematic representation of a generalized process layout of a preferred embodiment according to the present invention. The embodiment includes a supercritical reactor vessel with a material (5). A fluid from the fluid storage (1) is fed to the supercritical reactor vessel (5) at a controlled rate and under controlled conditions by means of the compressor (2) and a heat exchanger (3) for adjustment of fluid temperature. The compressor and heat exchanger forms the recirculation loop utilized for continuous control of the reactor conditions, particularly temperature and fluid composition. The fluid is withdrawn from the reactor through the separator (6), and recycled to the fluid storage (1). The alcohol produced in the reaction may be recollected in the separator (6) either during the reaction period, by circulating a purge stream of the supercritical fluid through the fluid storage (1).

20

The preparation of the fine particles may involve the following 3 steps:

1. Introduction of metal alkoxide or other pre-cursor into the supercritical reactor
2. introduction of the reaction promoter into the reactor
- 25 3. adjusting the reaction temperature and pressure to the desired level

Step 1, introduction of pre-cursor, may be performed by spraying a solution of the pre-cursor over the filling material of the reactor, while maintaining pressure and temperature in the reactor below the solubility limit of the pre-cursor, or by introducing a supercritical solution of the pre-cursor to the reactor, and reducing the solubility of the pre-cursor below the saturation point by appropriate change of pressure or temperature, and thereby causing a deposition of the pre-cursor on the filling material surface. The supercritical pre-cursor solution may be produced by introducing the pre-cursor solution through (4), while maintaining the supercritical solvent properties at an appropriate level by means of the recirculation loop.

35

In one embodiment supercritical solvent at appropriate conditions may be used to remove the solvent of the pre-cursor solution before entering into step 2.

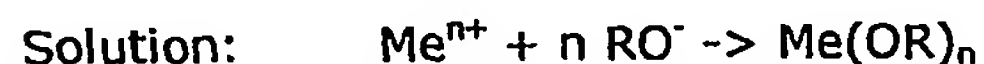
In step 2, the reaction promoter, preferably water, is introduced into the reactor. The introduction might take place by introducing the promoter directly into the reactor, or by introducing an at least partly saturated supercritical solution of the promoter. In a preferred embodiment the partly saturated supercritical solution of the promoter is  
 5 produced in an integrated recirculation loop, by introduction of the promoter through (4).

The adjustment of reaction conditions in step 3 may be performed by means of a recirculation loop for temperature control, and introduction or withdrawal of supercritical solvent to adjust the pressure, and thereby the solvent capacity.

10

It is understood that step 2 and 3 may be performed simultaneously in the same recirculation loop, or in any sequence, i.e. step 2 before step 3 or vice versa.

Fig. 8 shows schematic diagram of an in situ production alkoxide precursor according to  
 15 the present invention. The figure illustrates an electrochemical synthesis of said metal alkoxides being introduced at (4) in figure 7. The in-situ production may be provided by implemented using an anode (4) constructed from the metal to be transformed into the alkoxide, and a standard cathode (5). The electrodes are immersed into the alcohol solvent (3), and a suitable electric potential may be applied by means of a voltage generator (6).  
 20 The electrical conductivity of the alcohol solvent may be improved by addition of an organic salt or other suitable ionic species. The chemical reactions taking place may be:



In which Me denotes the metal, ROH the alcohol and  $\text{Me(OR)}_n$  a metal alkoxide. The overall  
 30 reaction is reduced to:



The formed hydrogen may be withdrawn through a vent (7), and the remaining metal  
 35 alkoxide solution may be withdrawn and introduced into said supercritical reactor through the outlet (2). The alcohol may be replenished through (1). Any ionic species added to modify the electrical conductivity of the solution may preferably be selected so as to be recollected with the excess alcohol, or purged out of the supercritical reactor during the pressurized state.

**ILLUSTRATIVE EXAMPLE 1:****Reactive production of fine particles as a nano-crystalline powder**

5

A preferred embodiment according to the present invention may be production of a fine particle material comprising nano-crystalline primary particles.

10 A generalized scheme for a batch process for production of a fine particles comprised by primary particles may involve the following consecutive steps:

- a. a dynamic pressurisation period,
- b. one or more reaction period(s) at elevated pressure,
- c. a depressurisation period.

15

A generalized schematic process diagram suitable for such production of fine particles in the form a powder comprising nanoscaled primary particles according to the present invention is disclosed in the figures 7-8, and the numbers below refers to these drawings.

20 The reactor (5) comprises one or more sections of a material for deposition of said primary particles. The material may be introduced into said vessel in the beginning of the method and withdrawn from said vessel at the end of the method, but preferably the material for powder production may be reused multiple times. The presence of the material within the vessel during the method may generate one or more of the following advantages in a least  
25 part time of the method:

- a. It serves as a flow distributor, thereby improving the distribution of said fluid(s) and/or substances being introduced and enabling a very precise control of the flow-, concentration-, pressure-, temperature- and density profiles within said  
30 vessel. A result from this improved distributing effect may be that more uniform and homogeneous primary particles are being produced.
- b. It provides a large number of nucleation sites so as to provide a high nucleation rate compared to the growth rate of the primary particles. Hereby, a seeding effect and/or a catalytic effect is/are introduced, thus ensuring a further improved  
35 control of the primary particles being formed.
- c. It serves as a collecting medium for effectively collecting the primary particles formed.
- d. The material with said primary particles deposited on the surface may comprise the final product.



The material may have any shape and comprise a number of different materials depending on the specific embodiment and application. The properties of the primary particles being formed by the method may at least partly be controlled by the selection of the material.

- 5 Typically the material is selected from one or more of the following criteria:
- a. It should be able to withstand the operating conditions such as the temperature during said method according to the present invention,
  - b. It should be able to adsorb at least one of the reactants on the surface.
  - 10 c. It should preferably have a high specific surface area.
  - d. It should allow ease of separation of said primary particles formed from said material, if the final product is not comprised by said primary particles on said surface of said material.
- 15 In the pressurisation period, the reactor vessel (5) is pressurised by adding one or more fluid(s) to the vessel to the vessel until the pressure in the reactor vessel exceeds the desired pressure for production of a powder comprising primary particles according to the present invention. The temperature in the reactor vessel may be controlled by conventional means such as controlling the inlet temperature of said fluid(s) to the reactor
- 20 vessel in a heat exchanger (3) before introducing said fluid(s) into said reactor vessel, and/or the temperature of the walls in said reactor vessel, e.g. by applying a jacketed reactor vessel with a heating or cooling fluid, electrical heating etc.

The rate of pressure increase may be constant, but many embodiments according to the

25 present invention involve a rate of pressure increase, which may vary according to a pre-selected schedule. Hence, many embodiments of the present invention involve controlling the rate of pressure increase to a pre-selected level e.g. the rate of pressure increase may typically be in the range 0,05-100 bar/min, such as in the range 0,1-20 bar/min and preferably in the range 0,1-15 bar/min, such as in the range 0,2-10 bar/min.

30 Particularly, the rate of pressure increase may be controlled to a pre-selected rate in at least part of the pressurisation period for economic reasons i.e. the pump or compressor size required may grow uneconomically big, and/or because the energy consumptions grow uneconomical and/or because the material in the vessel may not be able to withstand

35 the pressure rates being applied, and may lose its mechanical integrity.

In many embodiments according to the present invention, the rate of pressure increase may be controlled in the range 40-120 bars such as in the range 60-110 bars, and particularly in the range 65 to 110 bars. In a preferred embodiment according to the

present invention the rate of pressurisation in the interval 40 to 120 bars is at the most half of the maximum pressurisation rate outside this pressure range such as maximum one third of the maximum pressurisation rate this pressure range, and preferably at the most one fifth of the maximum rate of pressurisation outside this pressure range, such as  
5 maximum one tenth of the maximum rate of pressurisation outside this pressure range

One or more reactant(s) may be introduced into the reactor vessel before starting pressurisation period, but many preferred embodiments according to the present invention may involve introducing at least one of said reactant(s) during said pressurisation period  
10 or prior to said one or more reaction period(s). In embodiments, wherein said one or more reactants are being introduced before or during said pressurisation period, this may be performed by spraying a fluid containing said one or more reactant(s) over said material.

The pressurisation is performed by feeding one or more fluid(s) from a fluid storage (1) by  
15 one or more pump(s) and/or compressor(s) (2).

Said fluid storage may comprise a plurality of storage(s) for said fluid(s) so as to handle more than one fluid. The fluid(s) being fed to said reactor may comprise a gas or/ a liquid form of said fluid(s) or a combination of the two. For embodiments, wherein at least one of  
20 the fluid(s) is /are fed from said storage(s) in a liquid form, said liquid(s) is/are typically fed to an evaporator before introduction to said reaction vessel or mixing with another fluid and/or fluid mixture prior to introduction to said reactor vessel.

The pressure and temperature of fluid and/or fluid mixture in the reactor vessel after the  
25 pressurisation period are in many embodiments of the present invention maintained at a level, wherein at least one of said fluid(s) are in a supercritical state. The desired state of said supercritical fluid(s) prior to said reaction period may typically be selected so as to obtain a specific solubility of at least one reactant for said subsequent chemical reaction. Typically the pressure of said fluid(s) or fluid mixture in said reactor vessel may be in the  
30 range 85-500 bar such as in the range 100-300 bar prior to said reaction period.

As the pressurisation of said reactor vessel is achieved by introducing said fluid(s), and as the fluid generally is compressible, further compression takes place in the reactor vessel. The heat of compression may lead to a significant uncontrolled temperature increase in  
35 large vessel. If for example, carbon dioxide is compressed from 1 bar to 200 bar, the corresponding adiabatic temperature increase exceeds 100 C, which may lead to non-homogeneous reaction conditions within said reactor vessel, leading to undesirable large variations of said powder product to be produced in a method according to the present invention.

It is obvious to one skilled in the art, that the presence of a solid porous filling material in a significant part of the reactor vessel may hinder the extraction of heat through the vessel walls, as convective heat transfer is hindered, and the effect of this hindrance is  
5 proportional to the distance from the vessel centre to the vessel wall.

This may lead to undesirable large variations in the pressure- temperature- and density profiles within the vessel, which again may lead to reduced control of the primary particle formation reaction(s) in the subsequent reaction period(s) and/or may affect the  
10 mechanical integrity of said material being present in said one or more section(s) in said reactor vessel.

Hence, a preferred embodiment may involve controlling the pressure-, temperature- and/or density profiles within said reactor vessel during at least part time of said  
15 pressurisation period(s), by re-circulating at least part time of the method a part of the fluid contained in the vessel, the re-circulating comprising withdrawing from the reactor vessel at least part the fluid contained in the vessel, and withdrawing it to an external re-circulation loop for conditioning by e.g. extraction or addition of heat, and subsequently feeding the fluid to the vessel. The re-circulation during the pressurisation period enables  
20 very uniform and/or homogeneous reaction conditions with said reaction vessel prior to said reaction period(s). The re-circulation may further avoid excessively long pressurisation period(s) in order to achieve such preferred uniform reaction conditions, by enhancing the mass- and heat transfer rates in said reactor vessel. In a preferred embodiment according to the present invention the fluid in said re-circulation loop is in  
25 substantially, the same thermo-dynamical state as the fluid contained in said in the vessel, i.e. a gaseous state or a supercritical state.

Many embodiments according to the present invention involve one or more sol-gel reaction(s). In many of such embodiments a sequential or stepwise addition of said  
30 reactant(s) for said sol-gel reaction(s) is greatly preferred.

Often the least soluble reactant is introduced into the reactor vessel first. If for example the sol-gel reaction(s) involving the reaction between one or more alkoxide(s) and water, the alkoxide(s) may advantageously be introduced into the reactor vessel prior to  
35 introducing water. In a preferred embodiment said alkoxide(s) may be evenly distributed such as adsorbed or impregnated or coated on said the surface of said material being present in said one or more section(s) of said reactor vessel prior to introducing water.

The introduction of said alkoxide(s) may be introduced in a number of ways. The introduction may be performed by spraying a solution of said alkoxide(s) over said material, while maintaining pressure and temperature of said alkoxide(s) below the solubility limit(s) of said alkoxide(s). In another embodiment one or more supercritical  
5 fluid(s) with said alkoxide pre-cursor(s) dissolved and/or dispersed may be introduced into said vessel. Subsequently the density of said supercritical fluid(s) are decreased to a level below the solubility limit of said dissolved alkoxide(s) e.g. by an appropriate change of the pressure and/or temperature of said fluid, thereby causing a deposition of said alkoxide(s) on the surface of said material. In a still further embodiment the alkoxide(s) may be  
10 introduced into the vessel e.g. as dissolved in the corresponding alcohol, and said deposition of said alkoxide(s) obtained by introducing an antisolvent for said alcohol such as supercritical CO<sub>2</sub>, and thereby causing said alkoxide(s) to precipitate on the surface of said material. The supercritical CO<sub>2</sub> containing said dissolved alcohol may be withdrawn to the above described re-circulation loop for separation.

15

All the above described embodiments generally result in said alkoxide pre-cursor(s) being substantially uniformly distributed on the surface of said material being present in said one or more sections of said reactor vessel.

20 Subsequently another reactant(s) such as water may be introduced to said reactor vessel, preferably being dissolved in and/or mixed with a supercritical fluid. Said reactant(s) reacting with said well distributed alkoxide(s) on the surface of said material, and thereby causing primary particles to be formed on said surface of the material.

25 In some preferred embodiments according to the present invention, said sequential addition or stepwise addition may be repeated multiple times. The individual steps may comprise addition of the same reactants as added in the prior steps, or may comprise addition of new reactant(s).

30 The temperature and/or pressure may be maintained constant in some and/or all of the sequential steps, or may vary according to a pre-selected schedule.

Many preferred embodiments according to the present invention may further involve withdrawing from the reactor vessel at least part of the fluid contained in said vessel to a  
35 re-circulating loop for conditioning in at least part time of said one or more reaction period(s), or between said reaction period(s). In addition to controlling said pressure-, and/or temperature and/or density profiles according to a pre-selected schedule in some and/or all said reaction periods, said re-circulation may further control concentration

profiles e.g. by adding reactant(s) and/or extracting a reaction product(s) in at least part time of said reaction period(s) and/or between said reaction period(s).

In many embodiments according to the present invention the reactor vessel is  
5 depressurised, when the reaction period(s) is/are completed. The depressurisation may be performed in by applying similar suitable principles as described above for the pressurisation period(s). The fluid(s) may to a large extent be recovered during the depressurisation and recycled to the fluid storage(s).

- 10 The material comprising said primary particles on the surface produced according to the above description may comprise the final product. In such embodiments the product are extracted from the reactor vessel after the de-pressurisation period. Such products may further be subjected to an annealing process, such as a microwave annealing process.
- 15 In other embodiments the final fine particle product may constitute a powder comprised by said primary particles. Said primary particles may be easily separated from said material by introducing a vibrating effect and/or an acoustic effect such as ultra sound waves and/or by applying a back flushing effect, and/or a back shock effect and/or a back pulse effect.

20

In many embodiments of the present invention said powder product obtained by applying such means as described above comprises soft loosely bounded agglomerates, which are easily broken down to said primary particles by further processing. Said loosely bounded agglomerates may be easily collected by conventional separation means such as in a bag  
25 house filter or in a ceramic filter.

In a particularly preferred embodiment according to the present invention comprises separation of said powder product from said material within the vessel subsequent to said reaction period(s), thus allowing for semi-continuous or continuous operation. In such  
30 embodiment the method do not comprise said pressurisation and depressurisation periods described above.

In many embodiments according to the present invention powder product may have a high surface area such as measured by the BET method. The BET area of said powder product  
35 comprised by said primary may be at least 100 m<sup>2</sup>/g, such as at least 150 m<sup>2</sup>/g, preferably at least 200 m<sup>2</sup>/g such as at least 250 m<sup>2</sup>/g, and even more preferably at least 300 m<sup>2</sup>/g.



The present invention as illustrated in some preferred embodiments often allow very uniform and very homogeneous fine particle products to be produced at lower temperatures and/or shorter reaction times than prior art techniques. Typically a highly crystalline product may be obtained according to an embodiment of the present invention.

5 For clarity the crystallinity in this context shall preferably be interpreted as the weight ratio of a crystalline phase to the total weight of said fine particle material formed.

In many embodiments according to the present invention the crystallinity may be tuned to a specific level e.g. a crystallinity of more than 10 weight %, such as a crystallinity of more

10 30 weight %, preferably more than 50 weight % such as more than 70 weight %, and preferably more than 85 weight % such as substantially a crystalline material. In many embodiments according to the present invention the normally required post processing treatment steps such as drying and calcination steps are therefore eliminated or substantially reduced.

15

Due to the very precise and easy control of the process parameters in many embodiments according to the present invention, such embodiments are further considered to be suitable for large scale production.

## 20 **ILLUSTRATIVE EXAMPLE 2:**

An apparatus according to an embodiment of the present invention may include

- Reaction vessel assembly
- 25 • Dosing assembly for precursor and chemical reactor
- CO<sub>2</sub> recycle system
- Internal discharge assembly
- External filter and product collection assembly
- CO<sub>2</sub> storage assembly

30

The reaction vessel may be a vertical or a horizontal vessel. In a preferred embodiment a vertical vessel is used for facilities with a small production capacity and horizontal vessels are preferably used for facilities with large production capacity. In each case vessels may be arranged in parallel for optimal plant configuration as determined by a man skilled in

35 the art.

The reaction vessel may be equipped with one or more sections of high surface area material. The material is preferably arranged in a manner that allows easy cleaning and discharge from the high surface area material of the produced chemical reaction products.

Without limiting the scope of the invention the high surface area material may be hanging sheets of high surface area materials, hanging bags of high surface area materials or a honey come structured material. The reaction vessel further contains means for discharging the chemical reaction products from the high surface area material by using  
5 ultrasound, sonic horns, mechanical shaking, electrostatic discharge forces or any combination hereof.

The reaction vessel contains in the lower part means for collection and transport of the chemical reaction products. In a preferred embodiment a collection is performed using a  
10 mechanical conveyor that transports the formed particulate product to one end of the reaction vessel, where it is discharged into a pneumatic CO<sub>2</sub> transport system, which transport the products to an external filter and debugging system.

**CLAIMS**

1. A method of producing a fine particle material comprising
  - introducing one or more substances contained, such as dissolved and/or  
5 dispersed in one or more fluid(s) into a vessel by introducing said fluid(s) into the vessel, said vessel containing one or more section(s) comprising a material, at least one of the fluids being in a supercritical state before or after being introduced into said vessel,
  - causing and/or allowing, said substances to precipitate at least partly as primary  
10 particles on the surface of said material.
2. A method according to claim 1, wherein the average diameter of said nanoscaled primary particles is smaller than 100 nanometer such as smaller than 30 nanometer, preferably smaller than 20 nanometer, and even more preferable below 15 nanometer  
15 such as below 10 nanometer
3. A method according to any of the claims 1-2, wherein the standard deviation of the size distribution of said primary particles formed is less than 60 % of the average diameter of said primary particles, such as 40 % of the average diameter of said primary particles,  
20 and preferable less than 30 % of the average size of said primary particles, and even more preferable less than 20 % of the average size of said primary particles such as less than 15 % of the average size of said primary particles.
4. A method according to any of the claims 1-3, wherein the standard deviation of the  
25 size distribution of said primary particles formed is maximum 20 nanometer, such as maximum 10 nanometer, and preferably less than 5 nanometer, and even more preferably less than 3 nanometer.
5. A method according to any of the claims 1-4, wherein at least one of said fluid(s)  
30 being in a supercritical state is selected from the group consisting of carbon dioxide, alcohols such as methanol, ethanol, propanol, isopropanol, butanol, isobutanol, pentanol, hexanol, water, methane, ethane, propane, butane, pentane, hexane, cyclohexane, heptane, ammonia, sulfurhexafluoride, nitrous oxide, chlorotrifluoromethane, monofluoromethane, acetone, THF, acetic acid, citric acid, ethylene glycol, polyethylene  
35 glycol, N,N-dimethylaniline and mixtures thereof
6. A method according to claim 5, wherein one of the fluids is CO<sub>2</sub>.
7. A method according to claim 5, wherein one of the fluids is an organic solvent.

8. A method according to any of claim 5, wherein one of the fluids is water.
9. A method according to any of the preceding claims, wherein the pressure of at least one of said fluids is in the range 85-500 bar, preferably in the range 85-500 bar, such as in the range 100-300 bar.
10. A method according to any of the preceding claims, wherein the temperature in the vessel is maintained in the range 20-500 °C, such as 30-450 °C, and preferable in the range 35-200 °C, and more preferable in the range 40-150 °C.
11. A method according to any of the preceding claims, wherein said fluid further comprises at least one co-solvent.
12. A method according to claim 11, wherein the co-solvent is selected from the group consisting of alcohol(s), water, ethane, ethylene, propane, butane, pentane, hexane, heptane, ammonia, sulfurhexafluoride, nitrous oxide, chlorotrifluoromethane, monofluoromethane, methanol, ethanol, propanol, isopropanol, butanol, pentanol, hexanol, acetone, DMSO, THF, acetic acid, ethyleneglycol, polyethyleneglycol, N,N-dimethylaniline and mixtures thereof.
13. A method according to any of the preceding claims, wherein fluid further comprises one or more surfactants, said surfactants being preferably selected from the group consisting of hydrocarbons and fluorocarbons preferably having a hydrophilic/lipophilic balance value of less than 15, where the HLB value is determined according to the following formula:  $HLB = 7 + \text{sum}(\text{hydrophilic group numbers}) - \text{sum}(\text{lipophilic group numbers})$ .
14. A method according to any of the claims 1-13, wherein said precipitation is provided/caused by a change in the solubility of at least one of said substances.
15. A method according to claim 14, wherein said change in the solubility is provided/caused by an antisolvent present in the vessel.
16. A method according to claim 14, wherein said antisolvent is one of the fluids being introduced to the vessel.

17. A method according to any of the claims 14-16, wherein said change in solubility of at least one of said substances is provided/caused by expanding at least one of said fluids containing at least one of said substances into the vessel.
- 5 18. A method according to any of the claims 14-17, wherein said change in solubility is provided/caused by a change in the temperature of said fluid(s).
19. A method according to any of the preceding claims, wherein at least one of said substances undergoes a chemical reaction.
- 10 20. A method according to claim 19, wherein said chemical reaction(s) is/are a sol-gel reaction(s).
21. A method according to claim 20, wherein the maximum temperature in the vessel  
15 during said sol-gel reaction(s) is maintained below 400 C, such as below 300 C, preferably below 250 C such as below 200 C, and even more preferably below 150 C such as below 100 C.
22. A method according to claims 19-21, wherein at least one said substances  
20 undergoing a chemical reaction is a sulphate salt or a halide such as a chloride.
23. A method according to claims 19-21, wherein at least one said substances undergoing a chemical reaction is an alkoxide.
- 25 24. A method according to claim 23, wherein said alkoxide comprises a metal- or semi-metal alkoxide.
25. A method according to claim 24 comprising:
- 30 i. introducing into the vessel at least one of said reactant(s) and/or precursor(s) and/or initiator(s) and/or catalyst(s) for said chemical reaction
- ii. subsequently introducing into the vessel one or more substances  
35 dissolved and/or dispersed in at least one fluid.
- iii. and vice versa



26. A method according to claim 25, comprising multiple subsequent steps of

iv. introducing into the vessel at least one of said reactant(s) and/or precursor(s) and/or initiator(s) and/or catalyst(s) for said chemical reaction

v. subsequently introducing into the vessel one or more substances dissolved and/or dispersed in at least one fluid.

vi. and vice versa

27. A method according to claim 26, wherein said material in said one or more section(s) is capable of adsorbing at least one of said reactant(s) and/or precursor(s) and/or catalyst(s) on said material.

28. A method according to claim 27, wherein said reactant(s) and/or precursor(s) and/or catalyst(s) is/are adsorbed substantially in a monolayer of said material.

29. A method according to any of the claims 19-28, wherein the time for said chemical reaction(s) is less than 24 hours, such as less than 12 hours and preferable less than 8 hours, and even more preferable less than 4 hours.

30. A method according to any of the claims 19-28, wherein the time for said chemical reaction(s) is maximum 2 hours, such as maximum 1 hour, preferably less than 30 minutes and even more preferably less than 15 minutes.

31. A method according to any of the preceding claims, wherein said material present in said one more section(s) of said vessel, provides a distributing effect of said fluid(s) being introduced into said vessel.

32. A method according to claim 31, wherein the fraction of the total volume comprised by the said material in said one or more section(s) is up to 70 %, such as up to 50 %, preferably up to 30 %, and even more preferably up to 20 %.

33. A method according to any of the preceding claims, wherein said material present in said one or more section(s) of said vessel, comprises additional nucleation sites.

34. A method according to any of the preceding claims, wherein said material present in said one or more section(s) of said vessel, provides a seeding effect.

35. A method according to claim 33, wherein the number of nucleation sites is further increased by introducing an ultrasound and/or a vibrating surface effect.
- 5 36. A method according to any of the claims 33-35, wherein said material present in said one or more sections is a template for forming said primary particles into a specific shape, size, structure or phase.
37. A method according to claim 36, wherein said primary particles being produced is a  
10 least partly crystalline.
38. A method according to claim 35-37, wherein said seeding effect is at least partly provided by seed particles.
- 15 39. A method according to claim 38, wherein said seed particles is in a fluidised or suspended state.
40. A method according to the claims 38-39, wherein said seed particles comprises the same material as said primary particles.  
20
41. A method according to any of the claims 31-40, wherein said material present in said one or more sections comprises a porous structure such as a sheet, a spongeous or a grid structure.
- 25 42. A method according to any of the claims 31-40, wherein said material present in said one or more sections is a fibrous material.
43. A method according to claims 41-42, wherein said material present in said one or more sections has/have a hydrophilic surface.  
30
44. A method according to claims 41-42, wherein said material present in said one or more section has/have a hydrophobic surface.
45. A method according to any of the claims 41-44, wherein said material is a polymer  
35 material.
46. A method according to claim 45, wherein the material in the vessel is a polymer or elastomer such as selected from the group consisting of polyethylene, polypropylene, polystyrene, polyesters, polyethylene terephthalate, polyvinyl chloride, polyvinyl acetates,

polyoxymethylene, polyacryloamide, polycarbonate, polyamides, polyurethane, copolymers thereof, chlorinated products thereof, rubbers and chlorinated rubber, silicone rubbers, butadiene rubbers, styrene-butadiene-rubbers, isoprene polymers, vulcanised fluororubbers, silicone rubbers.

5

47. A method according to any of the claims 46, wherein said polymer material is polypropylene.

48. A method according to claim 45, wherein said fibrous material is an elastic  
10 material.

49. A method according to any of the claims 41-44, wherein said material is a ceramic material.

15 50. A method according to claim 49, wherein said material is a glass wool such as quartz wool.

51. A method according to claim 49, wherein said material is an aerogel.

20 52. A method according to claim 51 comprising the steps of  
- producing said aerogel material by a sol-gel reaction in an organic solvent  
- removing said organic solvent by extraction in supercritical CO<sub>2</sub>  
- drying at least partly said aerogel  
- forming said primary particles on the surface of said aerogel by a method  
25 according to any of the preceding claims.

53. A method according to any of the claims 41-52, wherein the specific surface area (m<sup>2</sup>/m<sup>3</sup>) of said material in said sections is above 500 m<sup>2</sup>/m<sup>3</sup>, such as 1000 m<sup>2</sup>/m<sup>3</sup>, such as above 10.000 m<sup>2</sup>/m<sup>3</sup>, and preferably above 50.000 m<sup>2</sup>/m<sup>3</sup> such as above 100.000 m<sup>2</sup>/m<sup>3</sup>.  
30

54. A method according to claim 53, wherein said material is a filtration medium.

55. A method according to any of the claims 53-54, wherein said material present in said one or more sections in the vessel further comprises at least one hollow tubular  
35 member comprising an inner and an outer surface, and having at least one end communicating with the outside of said vessel, at least part of said hollow tubular member(s) comprising a membrane, at least one of the fluids being introduced into the vessel through the membrane.

56. A method according to claim 55, wherein said hollow tubular member(s) is blocked in the one end being inside the vessel.
57. A method according to claim 56, wherein both ends of said hollow tubular member(s) are open and are communicating with the outside of said vessel.
58. A method according to claim 57, wherein said membrane contains at least one dense layer.
59. A method according to any of the preceding claims 55-58, wherein said membrane is a porous membrane, preferably being a membrane having pores within a range of 0,001-100 micron, such as a pores in the range 0,01-10 micron, and preferably being a membrane having pores within the range of 0,01- 0,1 micron.
60. A method according to any of the claims 55-59, wherein the inlet end of said hollow tubular member(s) is communicating with an inlet plenum, wherein an antisolvent is introduced, at least part of said antisolvent permeating said membrane surface and providing a controlled and uniform distribution of said antisolvent in said vessel.
61. A method according to any of the claims 59-60, wherein at least one of said fluids containing dissolved substances is introduced into the vessel through said hollow tubular member(s).
62. A method according to any of the claims 60 or 61, wherein at least two fluid streams are introduced into the vessel through two separate sets of hollow tubular member(s) each having an inlet plenum communicating with the outside of the vessel.
63. A method according to any of the claims 61-62, comprising contacting at least part of the inner surface of said hollow tubular member(s) with a first fluid, and contacting at least part of the outer surface(s) of said hollow tubular member(s) with a second fluid, a least part of said first fluid permeating through at least part the wall(s) of said hollow tubular member(s), and mixes with said second fluid substantially at the outer surface(s) of said hollow tubular member(s).
64. A method according to claim 63, wherein a micro-emulsion of said first fluid in said second fluid is formed.
65. A method according to any of the claims 61-64, wherein one of the fluids is water or contains water.

66. A method according to claim 65, wherein said water or water mixture contains one or more substances preferably being dissolved or dispersed therein.

5 67. A method according to any of the claims 61-66, wherein one of the fluids is CO<sub>2</sub> or an oil.

68. A method according to claim 67, wherein said fluid further comprises one or more surfactants.

10

69. A method according to any of the claims 64-68, wherein said microemulsion formed comprises a water core.

70. A method according to claim 69, wherein said water core comprises dissolved  
15 and/or dispersed substances.

71. A method according to the claims 69-70, wherein the diameter of said water core in said emulsions formed is in the range 0,5-15 times the diameter of the pores of the membrane part of said hollow tubular member(s) contacting said second fluid, such as in  
20 the range 1-10 times the diameter of the pores of the membrane part of said hollow tubular member(s) contacting said second fluid, and preferably in the range 2-4 times the diameter of the pores of the membrane part of said hollow tubular member(s) contacting said second fluid.

25 72. A method according to any of the claims 65-71, wherein said material present in said one or more sections, comprises two sets of hollow tubular member(s), both sets of said hollow tubular member(s) comprising a plenum and an outlet plenum communicating with the outside of said vessel, and wherein two different fluids can be contacted with the inner surface of said hollow tubular member(s), and where two different emulsion(s) of  
30 said fluids in said fluid contacting the outer surface of said hollow tubular member(s) are formed.

73. A method according to any of the claims 65-72, wherein a reaction is occurring in said microemulsion(s).

35

74. A method according to any of the claims 65-73, wherein said fluid containing said microemulsion(s) are used to dissolve and/or extract substances outside of said vessel.



75. A method according to any of the preceding claims, wherein said vessel is operating at a substantially constant pressure, such as operating continuously at a constant pressure during pre-selected time intervals.
- 5 76. A method according to any of the preceding claims, wherein said vessel is at a substantially constant pressure at more than one subsequent pre-selected pressure levels.
77. A method according to any the preceding claims, comprising re-circulating in at least part time of the method at least part of a fluid mixture present in the vessel, the re-
- 10 circulating comprising:
- withdrawing from the vessel at least a part of a fluid from the vessel and feeding it to a re-circulation loop and subsequently feeding the fluid back to the vessel.
- 15 79. A method according to claim 77, further comprising the step of controlling the temperature of the fluid in the re-circulation loop.
80. A method according to any of the claims 77-78, wherein heat is added and/or extracted from the fluid in the re-circulation loop.
- 20 81. A method according to any of the claims 77-79, wherein one or more reactant is added and/or extracted from the fluid in the re-circulation loop.
82. A method according to 80, wherein at least one of the reactants is an alcohol an
- 25 alkoxide and/or water.
83. A method according to claim 81, wherein a metal- or semi-metal alkoxid are produced in-situ in the process prior to being introduced to said vessel by said fluid.
- 30 84. A method according to any of the preceding claims, wherein the method comprises controlling the temperature- and/or pressure- and/or density- and/or concentration profiles within the vessel.
85. A method according to any of the preceding claims, wherein the temperature
- 35 profile within the vessel is controlled by controlling the temperature and flow rate of at least one fluid flowing inside said hollow tubular members.

86. A method according to any of the preceding claims, wherein said material present in said one or more sections in vessel with said precipitated primary particles thereon comprises the final product.

5 87. A method according to claim 85, wherein said product comprises primary particles deposited on a carrier film such as a tape cast.

88. A method according to claim 85, wherein said primary particles on said surface of said material constitutes a film or a coating.

10

89. A method according to claim 87, wherein said film or coating has one or more layer(s) each layer having a layer thickness of up to 1 micron, such as a layer thickness below 500 nanometer, preferable a layer thickness below 250 nanometer such as a layer thickness below 100 nanometer. Even more preferable the layer thickness of said film is  
15 below 50 nanometer, such as a layer thickness below 30 nanometer.

90. A method according to any of the claims 87-88, wherein said coating comprises multiple layers.

20 91. A method according to claim 89, where the layers comprises different materials.

92. A method according to any of the claims 87-90, wherein said product is further subjected to an annealing process.

25 93. A method according to claim 91, wherein said annealing is performed by microwaves.

94. A method according to claim 85, wherein said primary particles are deposited on the surface of said material in the form of small clusters of individual particles, and  
30 preferably as individual particles.

95. A method according to claim 93, wherein said clusters comprises up to 100 atoms , such as up to 50 atoms, and preferably less than 10 atoms, and even more preferably less than 5 atoms.

35

96. A method according to any of the claims 93-94, wherein said clusters or individual particles on the surface of said material are deposited as quantum dots.

97. A method according to any of the preceding claims, wherein said primary particles precipitated on said surface of the material present in said one or more section(s) are removed from said material as a powder.
- 5 98. A method according to claim 96, wherein said powder consists of weakly bounded agglomerates of primary particles.
99. A method according to claim 97, wherein said powder has weakly bounded agglomerates of a size of maximum 10 micron, such as up to 5 micron, and preferably up  
10 to 1 micron such as up to 500 nanometer.
100. A method according to any of the claims 96-98, wherein said powder is removed from said material by introducing a vibrating effect and/or an acoustic effect such as ultrasound waves and/or by back flushing and/or by applying an pressure pulse effect.  
15
101. A method according to claim 99, wherein said vibrating effect is generated by piezoelectric means.
102. A method according to claim 99, wherein said vibrating effect is generated by a  
20 magneto-restrictive means.
103. A method according to any of the claims 96-101, wherein said powder is removed from said material while said material is within the vessel.
- 25 104. A method according to claim 102, wherein said removal of said powder are performed according to a back flush or back pulse or a back chock technique.
105. A method according to any of the claims 102-103, wherein said removed powder are withdrawn from the vessel by flushing with a fluid or fluid mixture present in the  
30 vessel.
106. A method according to claim 104, wherein said fluid containing said formed powder is fed into a second vessel containing a liquid.
- 35 107. A method according to any of the claims 104-105, wherein said fluid containing said formed powder is expanded into said liquid thereby providing said formed powder material as a dispersion in said liquid.

108. A method according to claim 104, wherein said fluid containing said formed powder is fed to a bag filter or ceramic filter for separation of said formed powder material from the fluid.

5 109. A method according to claim 104, wherein said fluid containing said formed particulate material is fed to a membrane separation device.

110. A method according to claim 104, wherein said formed powder contained in said fluid is deposited on to a second solid in a second vessel.

10

111. A method according to claim 109, wherein said deposition is performed by spraying.

112. A method according to any of the preceding claims, wherein said precipitated  
15 primary particles on said material are exposed to one or more coating or encapsulation steps and/or reaction steps with one or more materials.

113. A method according to claim 111, wherein said coating or encapsulation step(s) is performed within the vessel.

20

114. A method according to any of the claims 111-112, wherein said coating or encapsulation step(s) is performed at least partly during harvesting/removing said particles from said material.

25 115. A method according any of the claims 111-113, wherein said coating or a further coating is performed in a second vessel.

116. A method according to any of the preceding claims, wherein said nanomaterial production process involves one of the following processes: RESS (rapid expansion of  
30 supercritical solutions), GAS (Gas Antisolvent), SAS (solvent Anti Solvent), SEDS (Solution Enhanced Dispersion by supercritical fluid), PCA (Precipitation with Compressed Antisolvent), PGSS (Precipitation from Gas-saturated Solutions) and variations thereof.

117. A method according to any of the preceding claims, wherein said primary particles  
35 comprises an electro-ceramic material.

118. A method according to any of the preceding claims, wherein said primary particles comprises a semi-conducting material.

119. A method according to any of the preceding claims, wherein said primary particles comprises a magnetic, ferromagnetic, paramagnetic, or superparamagnetic material.
120. A method according to any of the preceding claims, wherein said primary particles  
5 comprises a core-shell structure.
121. A method according to any of the preceding claims, wherein said core comprises a magnetic or ferro magnetic core.
- 10 122. A method according to any of the preceding claims, wherein said primary particles comprises a piezoelectrical material.
123. A method according to claim 121, wherein said piezoelectrical material comprises lead zirconate titanate,  $\text{Pb}(\text{Zr}_{0.52}\text{Ti}_{0.48})\text{O}_3$   
15
124. A method according to any of the preceding claims, wherein said primary particles comprises oxide(s), oxyhydroxide(s), hydroxide(s) such as metal oxide(s), semi-metal oxide(s), metal oxyhydroxide(s), semi-metal oxyhydroxide(s), metal hydroxide(s), semi-metal hydroxides and combinations thereof.  
20
125. A method according to any of the claims 116-123, wherein said oxides comprises oxides of one or more of the following elements: Al, Si, Ti, Zr, Zn, Fe, Ni, Co, Ce, Ge, Ba, Sr, W, La, Ta, Y, Mn, V, Bi, Sn, Te, Se, Ga, Be, Pb, Cr, Mg, Ca, Li, Ag, Au, Pt, Pd, Cd, Mo, Eu and combinations thereof.  
25
126. A method according to claim 124, wherein said oxides is selected among silica, alumina, zirconia, titania and combinations thereof.
127. A method according to any of the claims, wherein said primary particles comprises  
30 carbide(s) and/or nitride(s).
128. A method according to any of the claims 123-124, wherein said metal or semi-oxide(s) is/are precursor(s) for a thermoelectric material.
- 35 129. A method according to claim 127, wherein said precursor(s) is/are reduced in the process by adding a reducing agent to form a thermoelectric material.
130. A method according to claim 128, wherein said thermoelectric material formed comprises a clathrate.



131. A method according to any of the claims 127-129, said thermoelectric material comprises one or more of the elements: Ba, Bi, Te, Se, Zn, Sn, Sr, Ga, Ge, Pb, Cd, Sb, Ag, Si and combinations thereof.

5

132. A method according to claim 130, further comprising doping said primary particles of said thermoelectric material with metals and/or semi-metals.

133. A method according to claim 131, wherein said thermoelectrical material have a  
10 thermal conductivity at temperatures above 20 C of maximum 10 watts per meter Kelvin, such as maximum 5 watts per meter Kelvin, preferably maximum 3 watts per meter Kelvin, such as maximum 1,5 watts per meter Kelvin, and even more preferably a heat conductivity of maximum 1 watt per meter Kelvin.

15 134. A method according to any of the preceding claims, wherein said primary particles comprises one or more pharmaceutical and/or biological material(s).

135. A method according to any of the preceding claims, wherein said primary particles deposited on the surface of said surface provides an antibacterial effect of said surface.

20

136. A method according to any of the preceding claims, wherein said material comprises a high surface area material.

137. An apparatus comprising one or more of the means disclosed in any of the  
25 preceding claims and being adapted to carry out the method according to any of the preceding claims.

138. A product obtainable according to a method in any of the preceding claims.

30 139. A tape cast for tape casting, comprising primary particles deposited on a carrier film, wherein said primary particles have:

- 35
- a. an average diameter of less than 100 nanometer such as an average diameter of less than 30 nanometer, preferably an average diameter of smaller than 20 nanometer and even more preferable an average diameter below 15 nanometer such as below 10 nanometer.
  - b. a narrow size distribution around the average diameter characterized by having a maximum standard deviation of said distribution of maximum 20 nanometer, such as maximum 10 nanometer, and preferably less than 5

nanometer.

140. A tape cast according to claim 138, suitable for production of a ceramic material by tape casting.
- 5 141. A tape cast according to claim 139, wherein the sintering temperature is maximum 1100 C, such as maximum 1000 C, and preferable maximum 900 C, and even more preferably maximum 800 C, such as 700 C.
- 10 142. A tape cast according to claim 140, wherein the material being produced is a piezomotor produced from lead zirconate titanate tape cast.
143. An item having a hard nanocrystalline coating comprising primary particles of  $\text{Al}_2\text{O}_3$  and  $\text{ZrO}_2$  according to any of the preceding claims, wherein said coating has a hardness of  
15 at least 10 GPA, such as a hardness of at least 15 GPA, and preferably above 20 GPA, and even more preferably a hardness of at least 25 GPA.
144. An item having a hard nanocrystalline coating comprising primary particles of  $\text{Al}_2\text{O}_3$  and  $\text{ZrO}_2$  according to any of the preceding claims, wherein said coating has a scratch and  
20 wear resistance of at least 30 N, such as a scratch and wear resistance of at least 35 N, preferably a scratch and wear resistance of at least 40 N, and even more preferably a scratch resistance of at least 45 N.
145. An item having a hard nanocrystalline coating according to any of the claims 142-  
25 143, further comprising primary particles of ZnO.
146. A hard nanocrystalline coating according to any of the claims 142-144, wherein said coating is applied to a polymer or a glass material.
- 30 147. A mechanical part with a hard nanocrystalline coating according to any of the claims 142-145, wherein said coating is applied to the surface of said material.

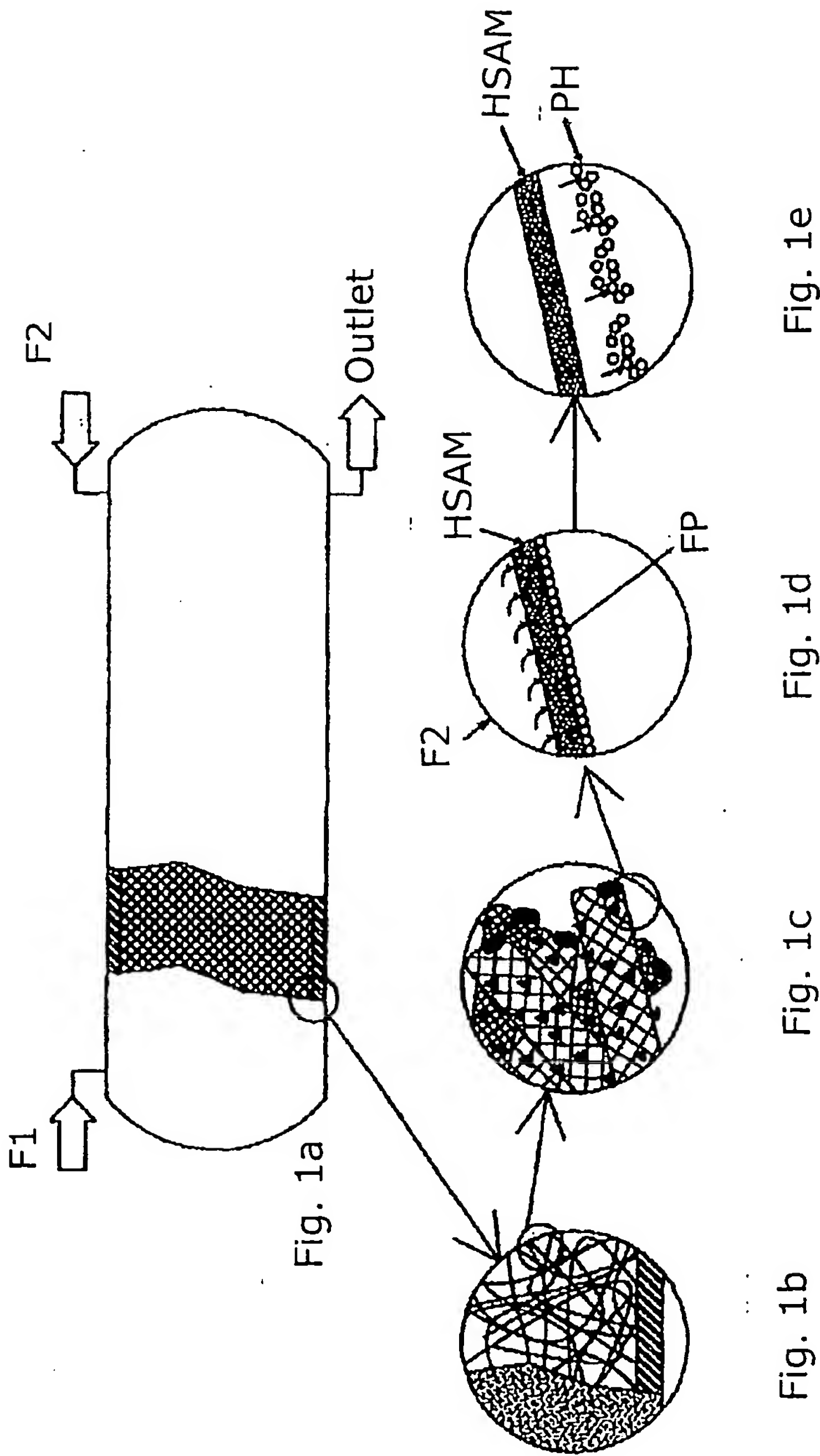


Fig. 1

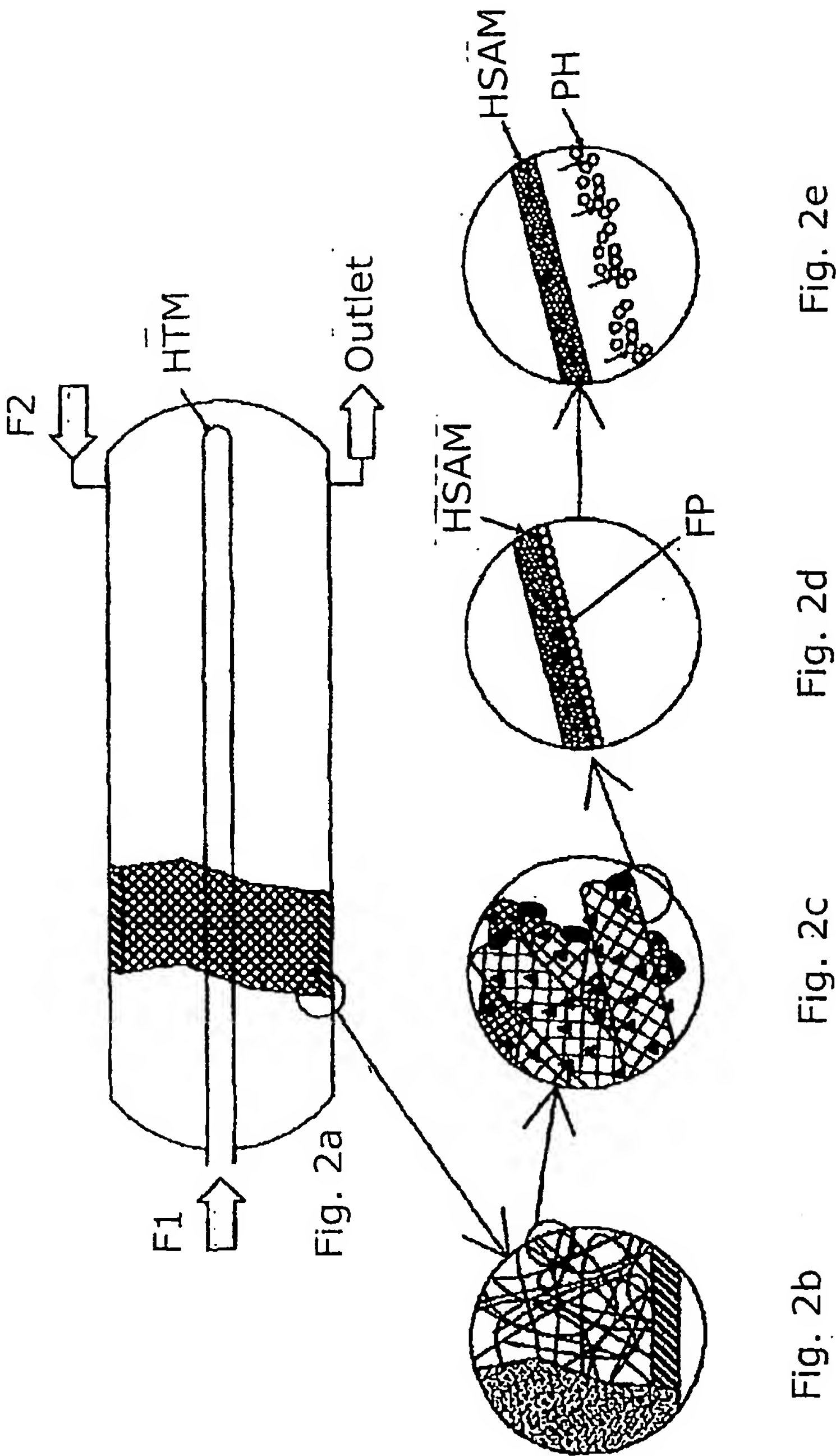


Fig. 2

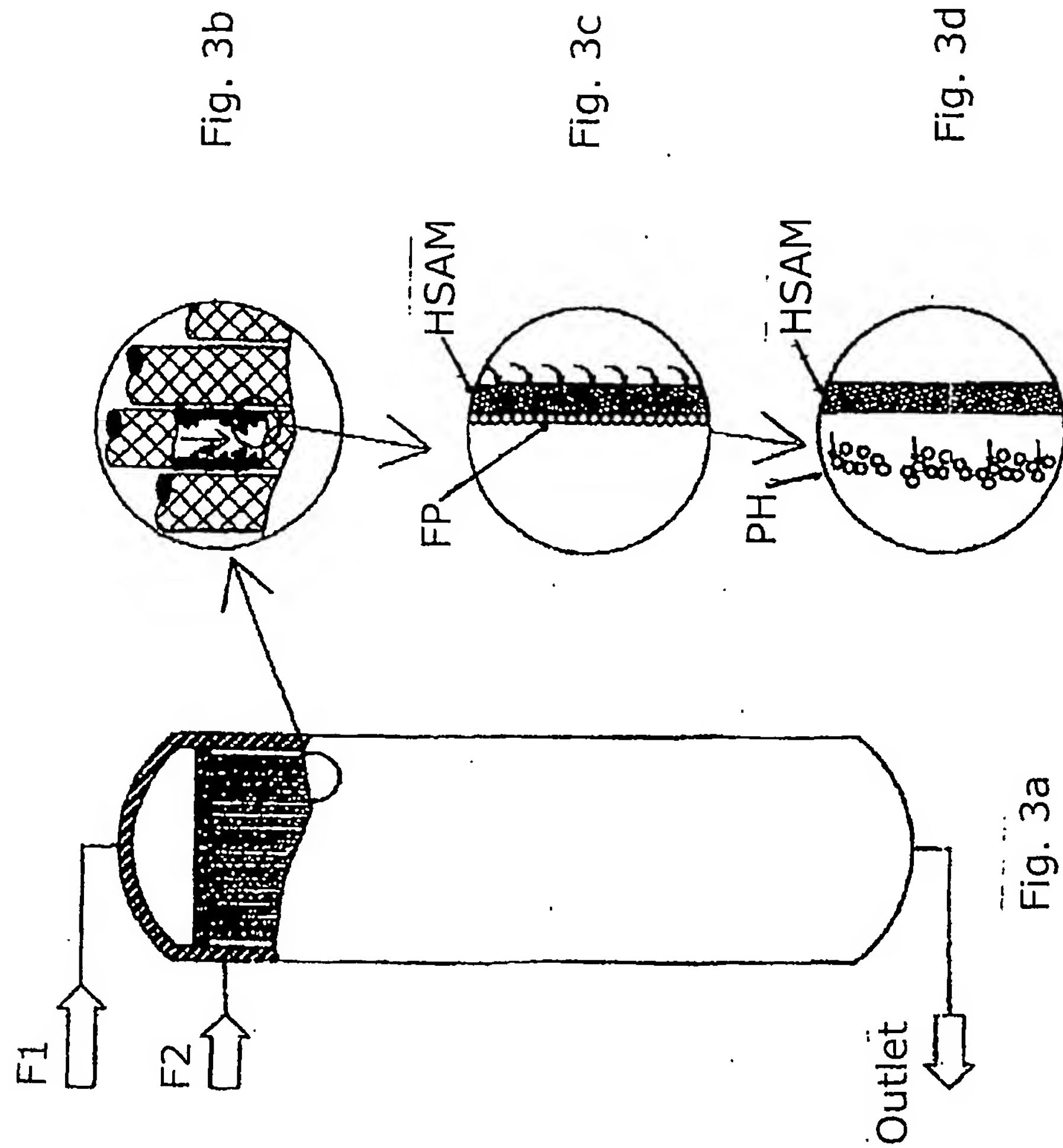


Fig. 3





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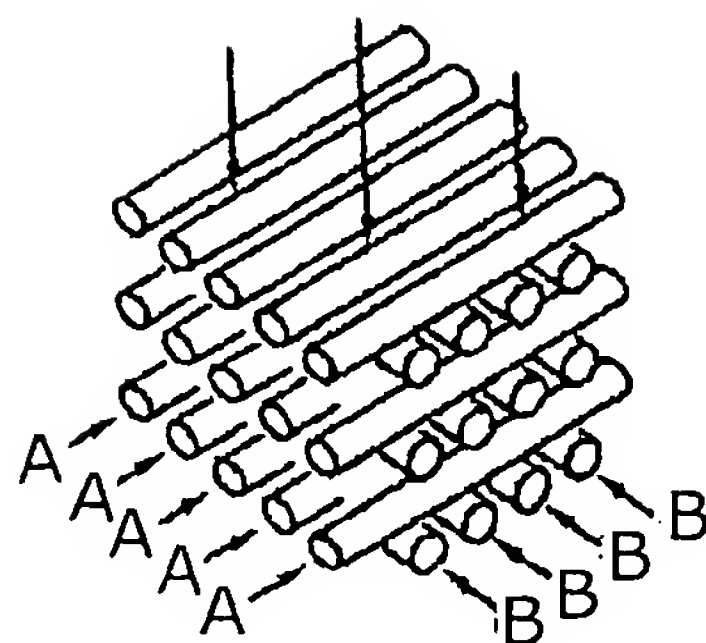


Fig. 5

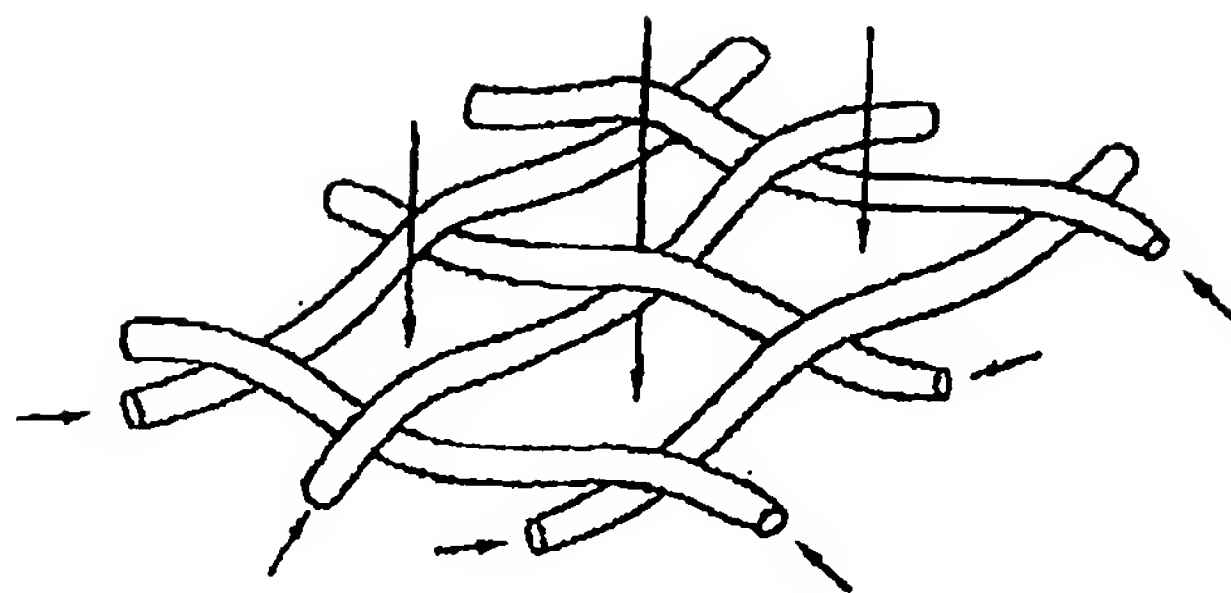


Fig. 6

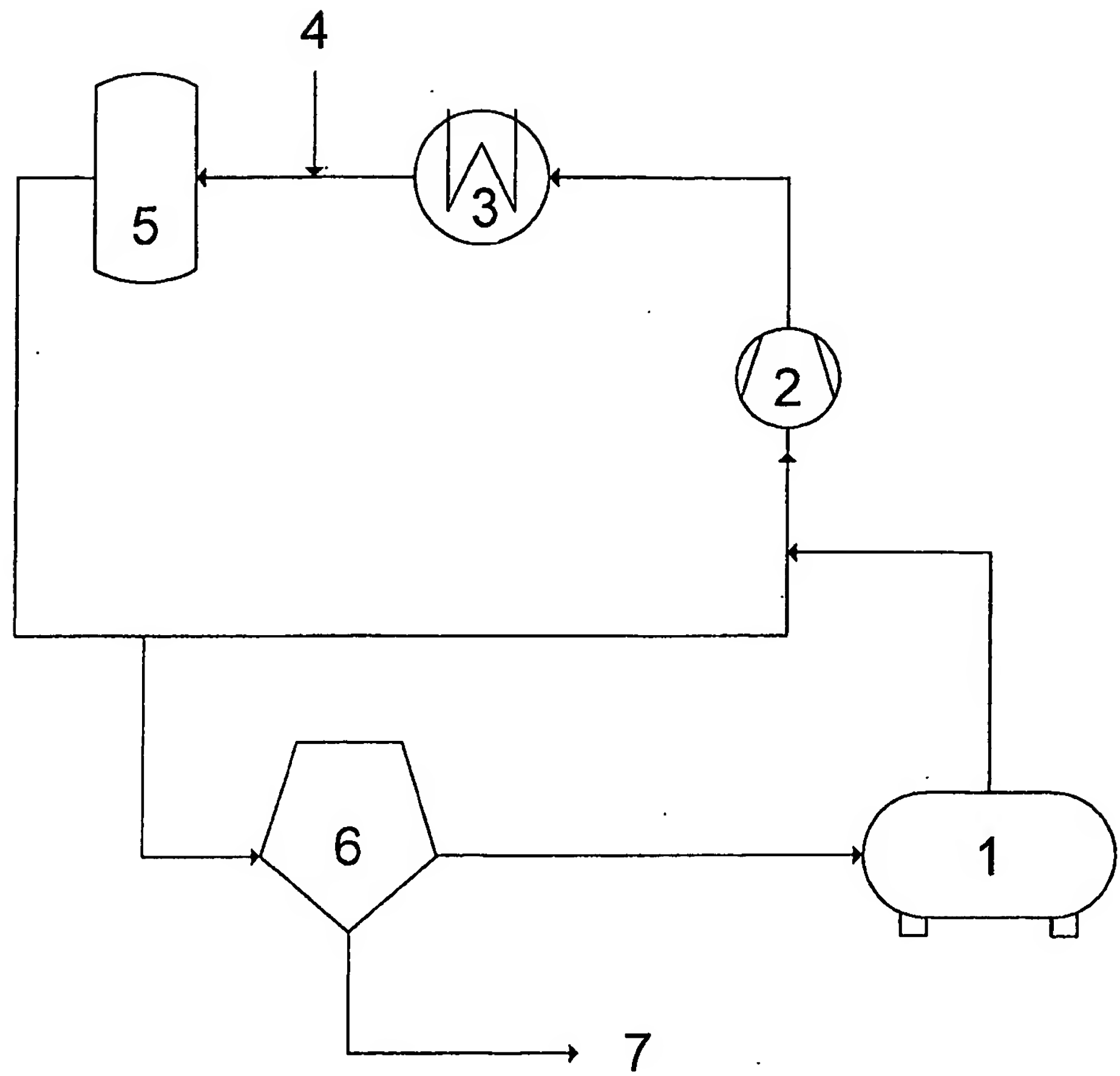
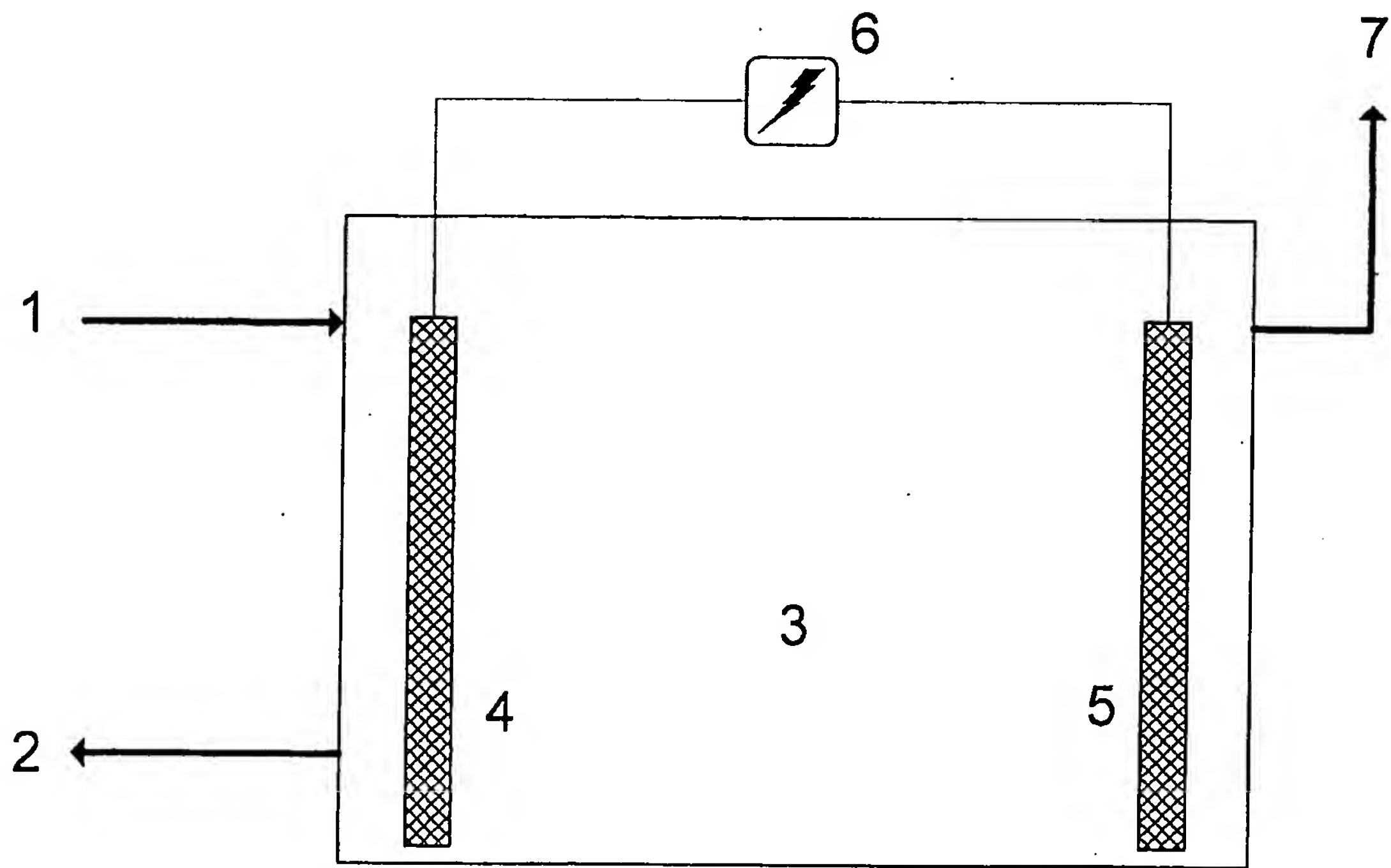


Fig. 7



**Fig. 8**  
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